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STRUCTURE FILE UPDATES: 30 JUL 2008 HIGHEST RN 1037244-07-7 DICTIONARY FILE UPDATES: 30 JUL 2008 HIGHEST RN 1037244-07-7

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http://www.cas.org/support/stngen/stndoc/properties.html

```
L2 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2008 ACS on STN
```

RN 13981-52-7 REGISTRY

ED Entered STN: 16 Nov 1984

CN Polonium, isotope of mass 210 (CA INDEX NAME)

OTHER NAMES:

CN 210Po

CN Po 210

CN Polonium-210

CN Radium F

DR 14809-83-7

MF Po

CI COM

LC STN Files: AGRICOLA, ANABSTR, BIOSIS, BIOTECHNO, CA, CAOLD, CAPLUS, CHEMLIST, CIN, CSNB, EMBASE, HSDB*, IFICDB, IFIPAT, IFIUDB, MEDLINE, MSDS-OHS, PIRA, PROMT, TOXCENTER, USPAT2, USPATFULL, USPATOLD (*File contains numerically searchable property data)

210_{Po}

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

3023 REFERENCES IN FILE CA (1907 TO DATE)

3 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

3024 REFERENCES IN FILE CAPLUS (1907 TO DATE)

42 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

- L3 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2008 ACS on STN
- RN 14255-04-0 REGISTRY
- ED Entered STN: 16 Nov 1984
- CN Lead, isotope of mass 210 (CA INDEX NAME)

OTHER NAMES:

- CN 210Pb
- CN Lead-210
- CN Pb 210
- CN Radium D
- MF Pk
- LC STN Files: AGRICOLA, ANABSTR, BIOSIS, BIOTECHNO, CA, CAOLD, CAPLUS, CHEMLIST, CIN, CSNB, EMBASE, IFICDB, IFIPAT, IFIUDB, PROMT, TOXCENTER, USPAT2, USPATFULL, USPATOLD

210_{Pb}

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 4375 REFERENCES IN FILE CA (1907 TO DATE)
 - 10 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
- 4376 REFERENCES IN FILE CAPLUS (1907 TO DATE)
 - 21 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

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FILE COVERS 1907 - 31 Jul 2008 VOL 149 ISS 5 FILE LAST UPDATED: 30 Jul 2008 (20080730/ED)

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'OBI' IS DEFAULT SEARCH FIELD FOR 'CAPLUS' FILE

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L2
             1 SEA FILE=REGISTRY ABB=ON 13981-52-7
L3
             1 SEA FILE=REGISTRY ABB=ON 14255-04-0
L4
         3024 SEA FILE=CAPLUS ABB=ON L2
L5
         4376 SEA FILE=CAPLUS ABB=ON L3
         1219 SEA FILE=CAPLUS ABB=ON L4 AND L5
L6
         2364 SEA FILE=CAPLUS ABB=ON RADIATION SOURCES/CT
L8
L9
             4 SEA FILE=CAPLUS ABB=ON L6 AND L8
             1 SEA FILE=REGISTRY ABB=ON 13981-52-7
L2
             1 SEA FILE=REGISTRY ABB=ON 14255-04-0
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L5
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L7
            28 SEA FILE=CAPLUS ABB=ON L2/P AND L3/P
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L10
          6038 SEA FILE=CAPLUS ABB=ON L10(L)(SOURCE#/OBI OR EMIT?/OBI)
L12
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L7
            28 SEA FILE=CAPLUS ABB=ON L2/P AND L3/P
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          248 SEA FILE=REGISTRY ABB=ON RADON?/CN
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L25
            6 SEA FILE=CAPLUS ABB=ON L7 AND L21
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L3
             1 SEA FILE=REGISTRY ABB=ON 14255-04-0
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L4
          4376 SEA FILE=CAPLUS ABB=ON L3
L5
L6
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           28 SEA FILE=CAPLUS ABB=ON L12 AND L6
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L20
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L21
      584884 SEA FILE=CAPLUS ABB=ON 71/SC,SX =NUCLEAR TECHNOLOGY
L24
             4 SEA FILE=CAPLUS ABB=ON L13 AND L21 AND L24
L26
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L3
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L5
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L7
           28 SEA FILE=CAPLUS ABB=ON L2/P AND L3/P
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         6038 SEA FILE=CAPLUS ABB=ON L10(L)(SOURCE#/OBI OR EMIT?/OBI)
L12
            28 SEA FILE=CAPLUS ABB=ON L12 AND L6
L13
L18
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L19 353488 SEA FILE=CAPLUS ABB=ON HYDROXIDE#/BI
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L3
              1 SEA FILE=REGISTRY ABB=ON 14255-04-0
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           4376 SEA FILE=CAPLUS ABB=ON L3
L5
L33
              5 SEA FILE=CAPLUS ABB=ON L4(L)PUR/RL
L34
              5 SEA FILE=CAPLUS ABB=ON L5(L)PUR/RL
              2 SEA FILE=CAPLUS ABB=ON L33 AND L34
L35
=> s 19,116,125,126,132,135
L82
            14 (L9 OR L16 OR L25 OR L26 OR L32 OR L35)
=> fil wpix; d que 181
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MOST RECENT THOMSON SCIENTIFIC UPDATE:
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                                                       <200848/DW>
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    20071130/UPIC and 20080401/UPIC.
    ECLA reclassifications to April and US national classifications to
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>>> Please note that the COPYRIGHT notification has changed <<<
'BI ABEX' IS DEFAULT SEARCH FIELD FOR 'WPIX' FILE
L70
             68 SEA FILE=WPIX ABB=ON (POLONIUM/BI, ABEX OR PO/BI, ABEX) (A) 210/BI
                ,ABEX OR 210PO/BI,ABEX OR PO210/BI,ABEX OR POLONIUM210/BI,ABEX
                OR 210POLONIUM/BI, ABEX
             68 SEA FILE=WPIX ABB=ON (LEAD/BI, ABEX OR PB/BI, ABEX) (A) 210/BI, ABE
L71
                X OR 210PB/BI, ABEX OR PB210/BI, ABEX OR 210LEAD/BI, ABEX OR
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LEAD210/BI, ABEX

L73	974355	SEA FILE=WPIX	ABB=ON	FILM#/BI,ABEX
L75	66573	SEA FILE=WPIX	ABB=ON	POLYCARBONATE#/BI,ABEX OR POLY CARBONATE#
		/BI,ABEX		
L76	146392	SEA FILE=WPIX	ABB=ON	HYDROXIDE#/BI,ABEX
L77	722352	SEA FILE=WPIX	ABB=ON	SEAL?/BI,ABEX
L78	1259	SEA FILE=WPIX	ABB=ON	RANDOM PULS?/BI,ABEX
L80	2652	SEA FILE=WPIX	ABB=ON	ALPHA/BI,ABEX(2A)(SOURCE/BI,ABEX OR
		EMIT?/BI,ABEX	OR PART	ICLE#/BI,ABEX)
L81	4	SEA FILE=WPIX	ABB=ON	L70 AND L71 AND (L73 OR L75 OR L76 OR
		L77 OR L78 OR	L80)	

=> fil PASCAL, BIOSIS, GEOREF, ENERGY, DISSABS, CONFSCI, INSPEC, EMBASE, COMPENDEX, SCISEARCH

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=> d que 149; d que 154; d que 156; d que 157; d que 163; d que 168

L37 7713 SEA (POLONIUM OR PO) (A) 210 OR 210PO OR PO210 L38 17263 SEA (LEAD OR PB) (A) 210 OR 210PB OR PB210 L42 47252 SEA POLYCARBONATE# OR POLY CARBONATE# L49 2 SEA L37 AND L38 AND L42

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L37
          7713 SEA (POLONIUM OR PO) (A) 210 OR 210PO OR PO210
L38
        17263 SEA (LEAD OR PB) (A) 210 OR 210PB OR PB210
L39
      2217612 SEA FILM#
L40
         68820 SEA RADON OR 222RADON OR RADON222
         98370 SEA ALPHA(2A)(SOURCE OR EMIT? OR PARTICLE#)
L41
L43
        190082 SEA HYDROXIDE#
L44
        226644 SEA SEAL?
L54
             9 SEA L37 AND L38 AND L39 AND ((L40 OR L41 OR L43 OR L44))
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L37
L38
          17263 SEA (LEAD OR PB) (A) 210 OR 210PB OR PB210
1.40
         68820 SEA RADON OR 222RADON OR RADON222
         98370 SEA ALPHA(2A)(SOURCE OR EMIT? OR PARTICLE#)
L41
L43
        190082 SEA HYDROXIDE#
L44
        226644 SEA SEAL?
             6 SEA L37 AND L38 AND L43 AND ((L40 OR L41 OR L44))
L56
L37
          7713 SEA (POLONIUM OR PO)(A) 210 OR 210PO OR PO210
L38
         17263 SEA (LEAD OR PB) (A) 210 OR 210PB OR PB210
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L40
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L41
        226644 SEA SEAL?
L44
L57
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         17263 SEA (LEAD OR PB) (A) 210 OR 210PB OR PB210
L38
L40
         68820 SEA RADON OR 222RADON OR RADON222
L58
         2389 SEA L37(2A) L38
          907 SEA L40(5A) COLLECT?
L61
             1 SEA L58 AND L61
L63
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L37
          17263 SEA (LEAD OR PB) (A) 210 OR 210PB OR PB210
L38
         68820 SEA RADON OR 222RADON OR RADON222
L40
L41
         98370 SEA ALPHA(2A)(SOURCE OR EMIT? OR PARTICLE#)
          2389 SEA L37(2A) L38
L66
          1616 SEA RANDOM PULS?
L68
              5 SEA L58 AND (L40 OR L41) AND L66
=> s 149,154,156,157,163,168
            23 (L49 OR L54 OR L56 OR L57 OR L63 OR L68)
=> dup rem 182,181,183
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PROCESSING COMPLETED FOR L82

PROCESSING COMPLETED FOR L81

PROCESSING COMPLETED FOR L83

L84 31 DUP REM L82 L81 L83 (10 DUPLICATES REMOVED)

ANSWERS '1-14' FROM FILE CAPLUS ANSWERS '15-16' FROM FILE WPIX ANSWERS '17-20' FROM FILE BIOSIS

ANSWERS '21-28' FROM FILE ENERGY ANSWER '29' FROM FILE EMBASE

ANSWER '30' FROM FILE COMPENDEX ANSWER '31' FROM FILE SCISEARCH

=> d ibib ab hitind 1-14; d iall abex tech 15-16; d iall 17-31; fil hom

L84 ANSWER 1 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN DUPLICATE 1

ACCESSION NUMBER: 2004:1156703 CAPLUS Full-text

DOCUMENT NUMBER: 142:44783

TITLE: Method for producing a sealed 210Pb-210Po

alpha source (alpha

particle emitter) and apparatus thereof

INVENTOR(S): Mitsuqashira, Hiroaki; Tsuyuzaki, Noriyoshi

PATENT ASSIGNEE(S): Japan

SOURCE: PCT Int. Appl., 21 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

	PAT	ENT	NO.			KIN.	D	DATE		-	APPL	ICAT	ION 1	NO.		D	ATE	
WO 2004114324 WO 2004114324			A2 A3		20041229 WO 2004-JP8407 20050224				20040609									
			AE, CN, GE, LR,	AG, CO, GH, LS,	CR, GM, LT,	AM, CU, HR, LU,	AT, CZ, HU, LV,		AZ, DK, IL, MD,	DM, IN, MG,	DZ, IS, MK,	EC, KE, MN,	EE, KG, MW,	EG, KP, MX,	ES, KR, MZ,	FI, KZ, NA,	GB, LC, NI,	GD, LK, NO,

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TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM,
             AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK,
             EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE,
             SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE,
             SN, TD, TG
                                            JP 2003-174296
     JP 2005010009
                          Α
                                20050113
                                                                   20030619
     EP 1634302
                         Α2
                                20060315
                                            EP 2004-736467
                                                                   20040609
         R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK
                                20060830
                                           CN 2004-80017060
     CN 1826663
                          Α
                                                                   20040609
     US 20070098606
                          Α1
                                20070503
                                            US 2005-560922
                                                                   20051215
                                                                A 20030619
PRIORITY APPLN. INFO.:
                                            JP 2003-174296
                                            WO 2004-JP8407
                                                                W 20040609
     A method is described for producing a sealed 210Pb-210Po \alpha source for a random
AΒ
     pulse generator. The method includes: collecting 210Pb-210Po with a 210Pb
     collector using Rn collection; precipitating the hydroxides of the collected
     210Pb-210Po and collecting the ppts. by a polycarbonate filter; dissolving the
     210Pb-210Po hydroxide precipitate to form a 210Pb-210Po radioactive thin film
     ; and sealing the 210Pb-210Po radioactive thin film for protection.
IC
    ICM G21G004-00
CC
    71-6 (Nuclear Technology)
ST
    radiation source lead polonium radon alpha particle
     emitter
ΙT
     Radiation sources
       (method for producing a sealed 210Pb-210Po alpha
       source)
ΙΤ
     Polycarbonates, uses
     Uranium ores
     RL: NUU (Other use, unclassified); USES (Uses)
        (method for producing a sealed 210Pb-210Po alpha
        source)
     13981-52-7P, Polonium-210, preparation 14255-04-0P,
ΤТ
     Lead-210, preparation
     RL: PUR (Purification or recovery); PREP (Preparation)
        (method for producing a sealed 210Pb-210Po alpha
        source)
     14859-67-7, Radon-222, uses
     RL: TEM (Technical or engineered material use); USES (Uses)
        (method for producing a sealed 210Pb-210Po alpha
        source)
L84 ANSWER 2 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN DUPLICATE 2
                         2003:91799 CAPLUS Full-text
ACCESSION NUMBER:
DOCUMENT NUMBER:
                         138:261770
TITLE:
                         Collection of emanating 222Rn for the preparation of a
                         210Pb-210Po alpha-source and the
                         building of a mobile random pulse and probability
                         generator utilizing alpha-counting technique
                         Hirose, N.; Tsuyuzaki, N.; Yamamot, H.; Mitsugashira,
AUTHOR(S):
                         T.; Hara, M.
CORPORATE SOURCE:
                         IWAKI Electronics Co., Ltd., Iwaki-city, Fukushima,
                         972-8322, Japan
                         Journal of Radioanalytical and Nuclear Chemistry
SOURCE:
                         (2003), 255(1), 207-210
                        CODEN: JRNCDM; ISSN: 0236-5731
                        Kluwer Academic Publishers
PUBLISHER:
                        Journal
DOCUMENT TYPE:
                        English
LANGUAGE:
```

AΒ A random pulse and probability generator (RPG) was developed using the detection technique of alpha-particles as the random signal source. collection technique for 222Rn emanated from natural U ore was examined for preparing highly pure 210Pb-210Po as an alpha source for RPG. The yield with a trap refrigerated by liquid N is >99% for 222Rn collection. CC 71-7 (Nuclear Technology) ST alpha source probability generator ΙT Distribution function (Poisson; in relation to alpha-source for random pulse and probability generator) TΤ Probability (emanating 222Rn in preparation of 210Pb-210Po alphasource for random pulse and probability generator) 12587-46-1, Alpha particle ТТ RL: PEP (Physical, engineering or chemical process); PYP (Physical process); PROC (Process) (decay detector; alpha-source for random pulse and probability generator) 14859-67-7, Radon-222, uses ΙΤ RL: NUU (Other use, unclassified); USES (Uses) (emanating 222Rn in preparation of 210Pb-210Po alphasource for random pulse and probability generator) 13981-52-7, Polonium-210, processes 14255-04-0, Lead-210, processes RL: PEP (Physical, engineering or chemical process); PYP (Physical process); PROC (Process) (emanating 222Rn in preparation of 210Pb-210Po alphasource for random pulse and probability generator) THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS REFERENCE COUNT: 5 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L84 ANSWER 3 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN DUPLICATE 3 ACCESSION NUMBER: 2000:551944 CAPLUS Full-text 133:156747 DOCUMENT NUMBER: Radiation sources for examination and/or calibration TITLE: of radiation detectors INVENTOR(S): Von Philipsborn, Henning PATENT ASSIGNEE(S): Germany

SOURCE: Ger. Offen., 8 pp.

CODEN: GWXXBX

DOCUMENT TYPE: Patent German LANGUAGE:

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 10002113	A1	20000810	DE 2000-10002113	20000119
PRIORITY APPLN. INFO.:			DE 1999-19902145	A1 19990120

AΒ A source for calibration comprises a radioelement-enriched carrier substrate. The radionuclide may be Pb-210 and/or Po-210. 1St a Rn-222-free substance is brought into a gas volume and then the Rn-222 concentration in the volume is increased. Then the gas contained in the volume is introduced into an organic solvent, whereby the Rn-222 contained in the gas dissolves in the solvent. The solvent is then aged over several days and poured over the carrier substrate through a screen end glass-fiber filter. Alternatively the carrier substrate can be also temporarily stored in the solvent. The process is lowcost and safe.

ICM G21G004-04 TC

ICS G01T001-16

```
CC 71-6 (Nuclear Technology)
IT Calibration
Radiation detectors
Radiation sources
(radiation sources for examination and/or calibration of radiation detectors)
IT 13981-52-7, Polonium-210, processes 14255-04-0,
Lead-210, processes 14859-67-7, Radon-222, processes
RL: PEP (Physical, engineering or chemical process); PROC (Process)
(radiation sources for examination and/or calibration of radiation detectors)
```

detectors)

4 ANSWER 4 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN

L84 ANSWER 4 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2007:71900 CAPLUS <u>Full-text</u> DOCUMENT NUMBER: 147:352120

TITLE: Estimate computing on source of nuclear and

radiological terrorism events

AUTHOR(S): Fu, Guang-zhi; Liu, Jun-feng; He, Bin; Zhang, Xi-xi CORPORATE SOURCE: The Second Artillery Engineering Institute, Xian of

Shaanxi Prov., 710025, Peop. Rep. China

SOURCE: He Dianzixue Yu Tance Jishu (2006), 26(6), 723-725,

716

CODEN: HDYUEC; ISSN: 0258-0934

PUBLISHER: Yuanzineng Chubanshe

DOCUMENT TYPE: Journal LANGUAGE: Chinese

- Once the nuclear and radiol. terrorism events have happened, and we do not know the nuclear kinds and amts., it should be very difficult for us to make a reasonable appraisement about the events, at the same time it might affect the protective and active, decision-making about meeting the emergency. One source anal. method was presented in this paper, and this method was used to explain an example of the source about the nuclear and radiol. events. This paper also pointed out that the mainly radiol. dangerous in the nuclear and radiol. terrorism events was long-lived α aerosol, and the mainly chemical dangerous was the very poisonous nuclide.
- CC 71-14 (Nuclear Technology)
- ST source nuclear radiol terrorism event alpha aerosol nuclide
- 7440-69-9, Bismuth (209), formation (nonpreparative) 13966-01-3, ΙT Thallium (210), formation (nonpreparative) 13966-27-3, Lead (206), formation (nonpreparative) 13966-29-5, Uranium (234), formation (nonpreparative) 13968-55-3, Uranium (233), formation (nonpreparative) 13981-14-1, Protactinium (233), formation (nonpreparative) 13981-52-7, Polonium (210), formation (nonpreparative) 13981-53-8, Radium (225), formation (nonpreparative) 13982-10-0, Plutonium (242), formation (nonpreparative) 13982-63-3, Radium (226), 13994-20-2, Neptunium (237), formation formation (nonpreparative) (nonpreparative) 14119-29-0, Lead (207), formation (nonpreparative) 14119-30-3, Lead (209), formation (nonpreparative) 14119-32-5, Plutonium 14119-33-6, Plutonium (240), formation (241), formation (nonpreparative) 14133-67-6, Thallium (207), formation (nonpreparative) (nonpreparative) 14255-04-0, Lead (210), formation (nonpreparative) 14265-85-1, Actinium (225), formation (nonpreparative) 14269-63-7, Thorium (230), 14331-79-4, Bismuth (210), formation formation (nonpreparative) (nonpreparative) 14331-85-2, Protactinium (231), formation (nonpreparative) 14596-10-2, Americium (241), formation (nonpreparative) 14733-03-0, Bismuth (214), formation (nonpreparative) 14835-02-0, Radon (219), formation (nonpreparative) 14859-67-7, Radon (222), formation (nonpreparative) 14932-40-2, Thorium (231), formation (nonpreparative) 15035-09-3, Thallium (206), formation (nonpreparative)

15065-10-8, Thorium (234), formation (nonpreparative) 15067-28-4, Lead (214), formation (nonpreparative) 15100-28-4, Protactinium (234), formation (nonpreparative) 15117-48-3, Plutonium (239), formation (nonpreparative) 15117-96-1, Uranium (235), formation (nonpreparative) 15229-37-5, Bismuth (211), formation (nonpreparative) 15422-74-9, Polonium (218), formation (nonpreparative) 15594-54-4, Thorium (229), formation (nonpreparative) 15623-45-7, Radium (223), formation (nonpreparative) 15623-47-9, Thorium (227), formation (nonpreparative) 15690-73-0, Thallium (209), formation (nonpreparative) 15706-52-2, Polonium (215), formation (nonpreparative) 15735-67-8, Polonium (214), formation (nonpreparative) 15735-83-8, Polonium (211), formation (nonpreparative) 15755-40-5, Astatine (218), formation (nonpreparative) 15756-41-9, Francium (221), formation (nonpreparative) 15756-57-7, 15756-98-6, Francium (223), Polonium (213), formation (nonpreparative) formation (nonpreparative) 15776-20-2, Bismuth (213), formation (nonpreparative) 15816-77-0, Lead (211), formation (nonpreparative) 17239-90-6, Astatine (217), formation (nonpreparative) 51696-22-1, Actinium (237), formation (nonpreparative) 51696-49-2, Thorium (237), formation (nonpreparative) RL: FMU (Formation, unclassified); POL (Pollutant); FORM (Formation, nonpreparative); OCCU (Occurrence)

(estimate computing on source of nuclear and radiol. terrorism events)

L84 ANSWER 5 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2006:256884 CAPLUS Full-text

DOCUMENT NUMBER: 144:365398

TITLE: Performance characteristics of sequential separation

and quantification of lead-210 and polonium-210 by ion

exchange chromatography and nuclear spectrometric

measurements

AUTHOR(S): El Afifi, E. M.; Borai, E. H.

Hot Laboratories and Waste Management Center (HLWMC), CORPORATE SOURCE:

Atomic Energy Authority, Cairo, Egypt

SOURCE: Journal of Environmental Quality (2006), 35(2),

568-574

CODEN: JEVQAA; ISSN: 0047-2425

PUBLISHER: American Society of Agronomy

Journal DOCUMENT TYPE: LANGUAGE: English

A selective separation and quant. determination procedure for 210Pb and 210Po AΒ in various environmental matrixes from different sources such as IAEA-326 soil, phosphate rocks (PR), and phosphogypsum (PG) was developed. The tested samples were digested sequentially using concentrated mineral acids (HF, HNO3) by a programmable high-pressure microwave digestion system. The sample solution was loaded onto a preconditioned ion exchange column (Sr-resin) for chromatog. separation Polonium-210 was eluted by 6 M HNO3 then spontaneously deposited onto polished silver disks to be measured using low-background alpha spectrometry. Lead-210 was sequentially eluted using 6 M HCl solution, precipitated as lead oxalate, dissolved in HNO3 solution, and mixed with scintillation cocktail to be measured by liquid scintillation counting (LSC). Performance of the developed procedure was tested using a reference soil (IAEA-326), with recommended isotope values, that was used as a quality control to assess separation and quantification efficiency (recovery %). The min. detectable activities of 210Pb and 210Po were found to be 24 and 0.28 Bq kg-1 for the measurements using LSC and alpha spectrometry, resp. The recoveries (%) of 210Pb and 210Po were found to be 80 and 60%, resp. To test the validity of the proposed LSC method, a comparative study was performed by measuring 210Pb activity concentration in test samples by nondestructive gamma-ray spectrometry. CC

8-1 (Radiation Biochemistry)

Section cross-reference(s): 19

IT 13981-52-7P, Polonium-210, analysis 14255-04-0P,
 Lead-210, analysis
 RL: ANT (Analyte); PUR (Purification or recovery); ANST
 (Analytical study); PREP (Preparation)
 (sequential separation and quantification of lead-210 and polonium-210 by ion exchange chromatog. and nuclear spectrometric measurements)

REFERENCE COUNT: 31 THERE ARE 31 CITED REFERENCES AVAILABLE FOR THIS

L84 ANSWER 6 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2003:535419 CAPLUS Full-text

DOCUMENT NUMBER: 139:313232

TITLE: Gamma coincidence study of 208Pb+350 MeV 64Ni

collisions

AUTHOR(S): Krolas, W.; Broda, R.; Fornal, B.; Pawlat, T.; Grawe,

H.; Maier, K. H.; Schramm, M.; Schubart, R.

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

CORPORATE SOURCE: H. Niewodniczanski Institute of Nuclear Physics,

Krakow, PL-31342, Pol.

SOURCE: Nuclear Physics A (2003), A724(3,4), 289-312

CODEN: NUPABL; ISSN: 0375-9474

PUBLISHER: Elsevier Science B.V.

DOCUMENT TYPE: Journal LANGUAGE: English

AB Products of 208Pb + 64Ni collisions at an energy 12% above the Coulomb barrier were studied in a γ -spectroscopy thick target experiment. The product yield distribution was established from the γ - γ coincidence anal. supplemented by target radioactivity measurements. Neutron evaporation from excited primary products was estimated to determine the pre-emission map of fragments. We discuss the transfer of protons and neutrons between the colliding ions in terms of the N/Z ratio equilibration.

CC 70-1 (Nuclear Phenomena)

7439-96-5P, Manganese 55, preparation 7440-03-1P, Niobium 93, ΙΤ 7440-25-7P, Tantalum 181, preparation 7440-38-2P, Arsenic preparation 75, preparation 7440-48-4P, Cobalt 59, preparation 7440-57-5P, Gold 197, preparation 7440-65-5P, Yttrium 89, preparation 7440-69-9P, Bismuth 209, preparation 10043-49-9P, Gold 198, preparation 10098-91-6P, Yttrium 90, preparation 10098-97-2P, Strontium 90, preparation 10198-40-0P, Cobalt 60, preparation 13965-98-5P, Krypton 83, preparation 13966-06-8P, Tin 113, preparation 13966-26-2P, Lead 204, preparation 13966-27-3P, Lead 206, preparation 13966-31-9P, Manganese 54, preparation 13967-66-3P, Iridium 191, preparation 13967-67-4P, Iridium 193, preparation 13967-71-0P, Zirconium 95, preparation 13967-73-2P, Strontium 85, preparation 13967-74-3P, Cerium 13967-76-5P, Niobium 95, preparation 141, preparation 13968-47-3P, Iron 58, preparation 13968-51-9P, Thallium 204, preparation 13968-53-1P, Ruthenium 103, preparation 13981-21-0P, Mercury 198, preparation 13981-25-4P, Copper 64, preparation 13981-27-6P, Zirconium 89, preparation 13981-29-8P, Terbium 160, preparation 13981-32-3P, Selenium 76, preparation 13981-37-8P, Nickel 63, preparation 13981-38-9P, Cobalt 58, preparation 13981-50-5P, Cobalt 57, preparation 13981-51-6P, Mercury 197, preparation 13981-52-7P, Polonium 210, preparation 13981-59-4P, Tin 117, preparation 13981-78-7P, Chromium 53, preparation 13981-80-1P, Nickel 60, preparation 13981-81-2P, Nickel 62, preparation 13981-83-4P, Cobalt 61, preparation 13981-84-5P, Cobalt 63, preparation 13982-00-8P, Tantalum 182, preparation 13982-08-6P, Ytterbium 170, preparation 13982-09-7P, Osmium 186, preparation 13982-12-2P, Rubidium 85, preparation 13982-13-3P, Rubidium 87, preparation 13982-14-4P, Strontium 86, preparation 13982-15-5P, Zirconium 90, preparation 13982-20-2P, Gold

193, preparation 13982-21-3P, Germanium 72, preparation 13982-22-4P, Gallium 72, preparation 13982-23-5P, Zinc 69, preparation 13982-24-6P, Iron 54, preparation 13982-30-4P, Cerium 139, preparation 13982-36-0P, Yttrium 88, preparation 13982-37-1P, Niobium 92, preparation 13982-38-2P, Bismuth 207, preparation 13982-39-3P, Zinc 65, preparation 13982-64-4P, Strontium 87, preparation 13982-78-0P, Mercury 203, preparation 13983-27-2P, Krypton 85, preparation 13994-19-9P, Xenon 127, preparation 14041-45-3P, Ytterbium 167, preparation 14041-50-0P, Ytterbium 171, preparation 14041-51-1P, Ytterbium 173, preparation 14041-52-2P, Ytterbium 172, preparation 14041-58-8P, Cadmium 114, preparation 14092-98-9P, Chromium 52, preparation 14093-02-8P, Iron 56, preparation 14093-09-5P, Hafnium 177, preparation 14093-11-9P, Hafnium 172, preparation 14093-12-0P, Lutetium 172, preparation 14107-52-9P, Thallium 197, preparation 14119-06-3P, Copper 65, preparation 14119-10-9P, Strontium 88, preparation 14119-12-1P, Zirconium 94, preparation 14119-15-4P, Molybdenum 99, preparation 14119-17-6P, Tin 120, preparation 14119-18-7P, Tin 122, preparation 14119-24-5P, Osmium 191, preparation 14119-28-9P, Lead 205, preparation 14119-29-0P, Lead 207, preparation 14119-30-3P, Lead 209, preparation 14133-67-6P, Thallium 207, preparation 14133-76-7P, Technetium 99, preparation 14145-42-7P, Bismuth 208, preparation 14158-27-1P, Strontium 89, preparation 14158-30-6P, Iodine 124, preparation 14158-35-1P, Iridium 194, preparation 14191-65-2P, Rubidium 89, preparation 14191-69-6P, Indium 116, preparation 14191-70-9P, Tin 116, preparation 14191-81-2P, Krypton 82, preparation 14191-82-3P, Krypton 86, preparation 14191-84-5P, Copper 63, preparation 14191-86-7P, Mercury 202, preparation 14191-87-8P, Mercury 199, preparation 14191-88-9P, Platinum 195, preparation 14234-24-3P, Yttrium 91, preparation 14255-04-0P, Lead 210, preparation 14265-71-5P, Selenium 75, preparation 14265-76-0P, Hafnium 179, preparation 14265-77-1P, Hafnium 178, preparation 14265-78-2P, Hafnium 180, preparation 14265-79-3P, Tungsten 180, preparation 14265-80-6P, Tungsten 182, preparation 14265-81-7P, Tungsten 183, preparation 14265-82-8P, Tungsten 184, preparation 14265-83-9P, Tungsten 186, preparation 14265-84-0P, Iridium 189, preparation 14269-78-4P, Ytterbium 169, preparation 14274-68-1P, Yttrium 87, preparation 14274-76-1P, Molybdenum 96, preparation 14274-79-4P, Osmium 190, preparation 14274-81-8P, Osmium 188, preparation 14276-53-0P, Copper 62, preparation 14280-37-6P, Bismuth 199, preparation 14280-38-7P, Bismuth 201, preparation 14280-48-9P, Thallium 203, preparation 14280-49-0P, Thallium 205, preparation 14304-78-0P, Arsenic 74, preparation 14304-79-1P, Tellurium 121, preparation 14304-80-4P, Tellurium 123, preparation 14304-97-3P, Chromium 54, preparation 14320-93-5P, Gold 195, preparation 14331-79-4P, Bismuth 210, preparation 14331-81-8P, Mercury 206, preparation 14331-90-9P, Bromine 84, preparation 14331-91-0P, Strontium 91, preparation 14331-93-2P, Zirconium 91, preparation 14333-38-1P, Bismuth 205, preparation 14336-70-0P, Nickel 59, preparation 14374-79-9P, Antimony 122, preparation 14374-81-3P, Germanium 71, preparation 14378-26-8P, Rhenium 188, preparation 14378-27-9P, Rhenium 190, preparation 14378-32-6P, Zinc 64, preparation 14378-33-7P, Zinc 66, preparation 14378-34-8P, Zinc 67, preparation 14378-35-9P, Zinc 68, preparation 14378-36-0P, Zinc 70, preparation 14378-53-1P, Rhodium 101, preparation 14380-59-7P, Bromine 81, preparation 14391-02-7P, Gallium 69, preparation 14391-03-8P, Gallium 71, preparation 14391-10-7P, Terbium 156, preparation 14391-11-8P, Gold 199, preparation 14391-27-6P, Tantalum 179, preparation 14391-28-7P, Rhenium 185, preparation 14391-29-8P, Rhenium 187, preparation 14391-61-8P, Bromine 80, preparation 14391-63-0P, Rubidium 82, preparation 14391-68-5P, Antimony 120, preparation 14391-73-2P, Copper 66, preparation

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14391-74-3P, Gallium 70, preparation 14391-76-5P, Silver 110,
preparation 14392-07-5P, Gadolinium 156, preparation 14392-15-5P,
Zirconium 92, preparation 14392-17-7P, Molybdenum 95, preparation
14392-19-9P, Molybdenum 97, preparation 14392-20-2P, Molybdenum 98,
preparation 14392-21-3P, Molybdenum 100, preparation 14452-48-3P,
Hafnium 176, preparation 14484-13-0P, Rhenium 183, preparation
14596-12-4P, Iron 59, preparation 14681-52-8P, Manganese 56, preparation
14681-54-0P, Selenium 80, preparation 14681-59-5P, Iron 55, preparation
14681-63-1P, Niobium 94, preparation 14681-65-3P, Niobium 90,
preparation 14681-72-2P, Selenium 77, preparation 14682-97-4P, Niobium
91, preparation 14683-00-2P, Molybdenum 94, preparation 14683-10-4P,
Antimony 124, preparation 14683-19-3P, Promethium 148, preparation
14683-24-0P, Gadolinium 154, preparation 14683-25-1P, Dysprosium 160,
preparation 14683-29-5P, Ytterbium 174, preparation 14683-32-0P,
Tungsten 179, preparation 14683-36-4P, Tantalum 183, preparation
14686-69-2P, Bromine 82, preparation 14687-25-3P, Lead 203, preparation
14687-40-2P, Germanium 75, preparation 14687-41-3P, Germanium 76,
preparation 14687-50-4P, Bismuth 202, preparation 14687-58-2P,
Selenium 82, preparation 14687-59-3P, Germanium 77, preparation
14687-60-6P, Selenium 83, preparation 14687-61-7P, Arsenic 77,
preparation 14687-62-8P, Bromine 83, preparation 14694-69-0P, Iridium
192, preparation 14762-69-7P, Iron 57, preparation 14809-52-0P,
Yttrium 85, preparation 14809-53-1P, Yttrium 86, preparation
14809-60-0P, Chromium 55, preparation 14809-62-2P, Cobalt 62,
preparation 14809-64-4P, Gallium 74, preparation 14809-66-6P, Arsenic
79, preparation 14809-68-8P, Krypton 87, preparation 14833-10-4P,
Vanadium 53, preparation 14833-16-0P, Selenium 78, preparation
14833-43-3P, Erbium 168, preparation 14833-49-9P, Nickel 65, preparation
14834-71-0P, Iridium 187, preparation 14834-72-1P, Promethium 143,
preparation 14834-83-4P, Ytterbium 166, preparation 14834-85-6P,
Dysprosium 162, preparation 14867-61-9P, Platinum 196, preparation
14900-10-8P, Erbium 164, preparation 14900-11-9P, Erbium 166,
preparation 14900-13-1P, Thulium 168, preparation 14900-21-1P, Hafnium
181, preparation
                 14903-04-9P, Bismuth 204, preparation 14913-25-8P,
Dysprosium 158, preparation 14913-50-9P, Thallium 208, preparation
14913-85-0P, Platinum 192, preparation 14913-89-4P, preparation
14914-16-0P, Gold 196, preparation 14914-52-4P, Zinc 71, preparation
14914-59-1P, Palladium 106, preparation 14914-60-4P, Ruthenium 100,
preparation 14914-61-5P, Ruthenium 101, preparation 14914-62-6P,
Ruthenium 102, preparation 14914-65-9P, Tin 118, preparation
14914-66-0P, Indium 117, preparation 14917-67-0P, Mercury 196,
preparation 14922-49-7P, Hafnium 174, preparation 14922-68-0P, Osmium
184, preparation 14922-70-4P, Platinum 188, preparation 14928-10-0P,
Nickel 61, preparation 14928-36-0P, Rubidium 88, preparation
14932-41-3P, Tungsten 185, preparation 14932-53-7P, Rubidium 86,
preparation 14981-91-0P, Iridium 190, preparation 14983-46-1P, Rhenium
184, preparation 14983-48-3P, Tungsten 187, preparation 14993-36-3P,
Osmium 182, preparation
RL: PNU (Preparation, unclassified); PREP (Preparation)
  (gamma coincidence study of 208Pb+350 MeV 64Ni collisions with light to
  heavy mass products)
14993-62-5P, Rhenium 180, preparation 14993-65-8P, Rhenium 181,
preparation 14993-91-0P, Krypton 84, preparation 14995-61-0P, Krypton
88, preparation 14998-63-1P, Rhenium 186, preparation 14998-72-2P, Tin 114, preparation 14998-96-0P, Platinum 194, preparation 14999-33-8P,
Manganese 53, preparation 15034-51-2P, Gallium 73, preparation
15034-58-9P, Germanium 73, preparation 15034-59-0P, Germanium 74,
preparation 15035-09-3P, Thallium 206, preparation 15047-33-3P,
Rhenium 179, preparation 15055-22-8P, Tantalum 178, preparation
15055-23-9P, Tungsten 178, preparation 15055-30-8P, Platinum 189,
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ΙT

preparation 15062-08-5P, Osmium 192, preparation 15064-65-0P, Thallium 201, preparation 15064-66-1P, Thallium 199, preparation 15064-97-8P, Mercury 194, preparation 15116-82-2P, Mercury 193, preparation 15125-00-5P, Rubidium 90, preparation 15125-53-8P, Tin 112, preparation 15185-19-0P, Mercury 201, preparation 15229-37-5P, Bismuth 211, preparation 15389-34-1P, Polonium 212, preparation 15411-62-8P, Ruthenium 99, preparation 15422-54-5P, Iron 61, preparation 15422-58-9P, Selenium 81, preparation 15422-59-0P, Arsenic 73, preparation 15575-20-9P, Arsenic 76, preparation 15678-91-8P, Krypton 81, preparation 15690-73-0P, Thallium 209, preparation 15691-06-2P, Zirconium 96, preparation 15700-41-1P, Niobium 98, preparation 15700-42-2P, Niobium 100, preparation 15700-83-1P, Bromine 85, preparation 15701-21-0P, Tantalum 184, preparation 15706-36-2P, Platinum 191, preparation 15706-38-4P, Polonium 204, preparation 15715-01-2P, Astatine 213, preparation 15715-06-7P, Mercury 192, preparation 15720-36-2P, Cobalt 64, preparation 15720-38-4P, Copper 68, preparation 15720-40-8P, Copper 69, preparation 15720-45-3P, Polonium 207, preparation 15720-55-5P, Thallium 200, preparation 15720-57-7P, Thallium 202, preparation 15735-68-9P, Platinum 190, preparation 15735-70-3P, Platinum 193, preparation 15735-74-7P, Platinum 197, preparation 15735-81-6P, Polonium 209, preparation 15735-83-8P, Polonium 211, preparation 15735-86-1P, Polonium 206, preparation 15735-87-2P, Polonium 208, preparation 15741-32-9P, Lutetium 170, preparation 15741-33-0P, Manganese 57, preparation 15743-50-7P, Thallium 198, preparation 15743-54-1P, Ytterbium 168, 15743-55-2P, Zinc 72, preparation 15749-40-3P, Titanium preparation 52, preparation 15749-46-9P, Tungsten 181, preparation 15749-58-3P. Palladium 108, preparation 15750-13-7P, Hafnium 175, preparation 15751-77-6P, Zirconium 93, preparation 15752-22-4P, Iridium 188, preparation 15752-27-9P, Lutetium 171, preparation 15752-86-0P, Lead 202, preparation 15755-35-8P, Arsenic 78, preparation 15755-36-9P, Astatine 212, preparation 15755-38-1P, Astatine 209, preparation 15755-39-2P, Astatine 211, preparation 15756-10-2P, Mercury 200, preparation 15756-14-6P, Mercury 204, preparation 15756-15-7P, Mercury 15756-45-3P, Gold 192, preparation 15756-63-5P, 195, preparation Platinum 198, preparation 15756-69-1P, Polonium 202, preparation 15756-83-9P, Germanium 78, preparation 15756-89-5P, Gold 194, preparation 15756-97-5P, Francium 212, preparation 15757-14-9P, Gallium 68, preparation 15757-23-0P, Hafnium 173, preparation 15757-86-5P, Copper 67, preparation 15758-18-6P, Chromium 56, preparation 15758-45-9P, Selenium 79, preparation 15758-49-3P, Strontium 84, preparation 15759-29-2P, Tantalum 180, preparation 15761-06-5P, Osmium 189, preparation 15765-69-2P, Radon 211, preparation 18765-70-8P, Radon 212, preparation 15765-78-3P, Rhenium 189, preparation 15765-86-3P, Rubidium 84, preparation 15766-01-5P, Ruthenium 104, preparation 15766-16-2P, Nickel 67, preparation 15766-33-3P, Nickel 66, preparation 15766-50-4P, Osmium 185, preparation 15766-52-6P, Osmium 187, preparation 15776-19-9P, Bismuth 206, preparation 15816-77-0P, Lead 211, preparation 15816-99-6P, Iridium 195, preparation 15832-32-3P, Niobium 96, preparation 15832-38-9P, Gold 190, preparation 15832-41-4P, Indium 118, preparation 15840-05-8P, Erbium 162, preparation 16394-57-3P, Gold 191, preparation 16468-57-8P, Germanium 79, preparation 16645-99-1P, Lead 200, preparation 16646-00-7P, Lead 198, preparation 16729-68-3P, Mercury 205, preparation 16729-74-1P, Polonium 203, preparation 16729-76-3P, Polonium 205, preparation 17056-36-9P, Rubidium 83, preparation 17058-33-2P, Radon 209, preparation 17239-85-9P, Bismuth 200, preparation 17239-87-1P, Lead 201, preparation 17620-09-6P, Francium 213, preparation 17638-03-8P, Radium 214, preparation 18390-45-9P, Astatine 208, preparation 18476-92-1P,

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Manganese 58, preparation 18496-04-3P, Niobium 97, preparation
18624-12-9P, Cobalt 66, preparation 18724-77-1P, Thallium 196,
preparation 18830-37-0P, Astatine 210, preparation
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Gallium 75, preparation 21410-52-6P, Gallium 76, preparation 21459-51-8P, Astatine 206, preparation 21459-71-2P, Rhenium 182,
preparation
              21459-72-3P, Osmium 183, preparation
                                                     22453-47-0P, Platinum
187, preparation 22453-70-9P, Niobium 99, preparation
                                                           24383-94-6P,
Bismuth 203, preparation 24447-13-0P, Iridium 186, preparation
25731-76-4P, Yttrium 84, preparation 26110-67-8P, Krypton 80,
              26683-69-2P, Thallium 195, preparation 27485-99-0P, Lead
preparation
197, preparation 27486-00-6P, Lead 199, preparation 27742-26-3P,
Iridium 184, preparation 28637-43-6P, Vanadium 54, preparation
29054-43-1P, Iridium 185, preparation 29136-29-6P, Radon 214,
preparation 29675-20-5P, Iron 62, preparation 29675-21-6P, Nickel 68,
preparation 29675-28-3P, Copper 70, preparation 29675-32-9P, Titanium
53, preparation 29675-34-1P, Nickel 69, preparation 29675-35-2P, Zinc
73, preparation 29675-38-5P, Cobalt 65, preparation 30017-28-8P,
Copper 71, preparation 32020-21-6P, Iron 60, preparation 32025-57-3P, Radon 213, preparation 33233-20-4P, Radon
210, preparation 33690-55-0P, Arsenic 80, preparation
                                                            36819-19-9P,
Vanadium 55, preparation 36819-21-3P, Chromium 57, preparation
36819-22-4P, Manganese 59, preparation 42250-70-4P, Chromium 58,
              42250-73-7P, Manganese 60, preparation 52813-79-3P,
preparation
Manganese 61, preparation 58831-77-9P, Iron 64, preparation
58831-78-0P, Iron 63, preparation 72062-02-3P, Manganese 62, preparation
RL: PNU (Preparation, unclassified); PREP (Preparation)
   (gamma coincidence study of 208Pb+350 MeV 64Ni collisions with light to
   heavy mass products)
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REFERENCE COUNT: 23 THERE ARE 23 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L84 ANSWER 7 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2004:56675 CAPLUS <u>Full-text</u>

DOCUMENT NUMBER: 141:29931

TITLE: Preparation of low level sealed 210Pb source

for random pulse generator

AUTHOR(S): Mitsugashira, T.; Hara, M.; Tsuyuzaki, N.

CORPORATE SOURCE: Institute for Materials Research, Tohoku University,

Japan

SOURCE: KEK Proceedings (2003), 2003-11(Proceedings of the

Fourth Workshop on Environmental Radioactivity, 2003),

263-268

CODEN: KEPREW

PUBLISHER: High Energy Accelerator Research Organization

DOCUMENT TYPE: Journal LANGUAGE: Japanese

Two methods for sealed 210Pb-210Po source preparation, direct deposition from isoPr alc. solution (IPA solution) of 210Pb-210Po nitrate (D-IPA method) and the 210Pb-210Po hydroxides precipitation (PPT method), were exptl. examined In the former D-IPA method, an aliquot of IPA solution of 210Pb-210Po nitrate was directly dropped in a sealed cap for PPD and dried by heating. Then, a polycarbonate (PC) solution of 1/1 mixture of CH2Cl2 and dichloroethane was dropped on the source to make a thin (.apprx.0.1 mg/cm2) film for radioactivity sealed . In the PPT method, 210Pb-210Po hydroxide was filtered on a PC membrane filter (Nuclipore 0.1 μ m) and the membrane filter was dissolved in a 1/1 mixture of CH2Cl2 and dichloroethane. The sealed 210Pb-210Po sources were prepared directly by dropping an aliquot of the solution into the PPD cap followed by its evaporation. The sealed sources thus prepared were subjected to 1 m height drop test, air blowing test, and H2O immersion

test. No radioactive contaminants were coming off from the sealed source through these tests. CC 71-6 (Nuclear Technology) ST alpha particle source polonium sealing polycarbonate membrane ΙT Membrane filters (for preparation of low level sealed 210Pb source for random pulse generator) ΙT Polycarbonates, uses RL: NUU (Other use, unclassified); USES (Uses) (for preparation of low level sealed 210Pb source for random pulse generator) Radiation sources ΙT Sealing (preparation of low level sealed 210Pb source for random pulse 67-63-0, Isopropyl alcohol, uses 75-09-2, Dichloromethane, uses ΙT 1300-21-6, Dichloroethane 12027-17-7, Polonium hydroxide (PO(OH)4 127795-35-1, Polonium nitrate RL: NUU (Other use, unclassified); USES (Uses) (for preparation of low level sealed 210Pb source for random pulse generator) 12587-46-1, Alpha ray ΤТ RL: FMU (Formation, unclassified); FORM (Formation, nonpreparative) (preparation of low level sealed a source for random pulse generator) ΙΤ 13981-52-7, Polonium-210, uses 14255-04-0, Lead-210, RL: TEM (Technical or engineered material use); USES (Uses) (preparation of low level sealed 210Pb source for random pulse generator) L84 ANSWER 8 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN 2002:893088 CAPLUS Full-text ACCESSION NUMBER: DOCUMENT NUMBER: 138:261764 TITLE: In vitro cell irradiation systems based on 210Po alpha source: construction and characterization Szabo, J.; Feher, I.; Palfalvi, J.; Balashazy, I.; AUTHOR(S): Dam, A. M.; Polonyi, I.; Bogdandi, E. N. CORPORATE SOURCE: KFKI Atomic Energy Research Institute, Budapest, H-1525, Hung. Radiation Measurements (2002), 35(6), 575-578 SOURCE: CODEN: RMEAEP; ISSN: 1350-4487 PUBLISHER: Elsevier Science Ltd. DOCUMENT TYPE: Journal LANGUAGE: English One way of studying the risk to human health of low-level radiation exposure AB is to make biol. expts. on living cell cultures. Two 210Po α -particle emitting devices, with 0.5 and $100~\mathrm{MBq}$ activity, were designed and constructed to perform such expts. irradiating monolayers of cells. Ests. of dose rate at the cell surface were obtained from measurements by a PIPS α -particle spectrometer and from calcns. by the SRIM 2000, Monte Carlo charged particle transport code. Particle fluence area distributions were measured by solid state nuclear track detectors. The design and dosimetric characterization of the devices are discussed. 71-7 (Nuclear Technology) CC Section cross-reference(s): 8 Animal tissue culture ΤТ

Dosimeters

Radiation sources

(210Po α -particle emitting devices: design and dosimetric characterization for in vitro radiobiol. studies)

13981-52-7, 210Po, uses

RL: BUU (Biological use, unclassified); DEV (Device component use); BIOL (Biological study); USES (Uses)

(210Po $\alpha\text{-particle}$ emitting devices: design and dosimetric characterization for in vitro radiobiol. studies)

7440-69-9, Bismuth-209, uses 14255-04-0, 210Pb, uses ΙT 14331-79-4, 210Bi, uses

RL: PEP (Physical, engineering or chemical process); PYP (Physical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses)

(210Po α -particle emitting devices: design and dosimetric characterization for in vitro radiobiol. studies)

THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS REFERENCE COUNT: 6 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L84 ANSWER 9 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 1997:743658 CAPLUS Full-text

DOCUMENT NUMBER: 128:67547

ORIGINAL REFERENCE NO.: 128:13107a,13110a

TITLE: A new method for evaluating annual alpha and beta dose

rates in different ceramic samples by using solid

state nuclear track detectors

Misdag, M. A.; Fahde, K.; Erramli, H.; Mikdad, A. AUTHOR(S): CORPORATE SOURCE: Nuclear Physics and Techniques Laboratory, Faculty of

Sciences Semlalia, University Cadi Ayyad, Marrakech,

Morocco

SOURCE: Radiation Physics and Chemistry (1997), 50(3), 293-297

CODEN: RPCHDM; ISSN: 0969-806X

PUBLISHER: Elsevier Science Ltd.

DOCUMENT TYPE: Journal LANGUAGE: English

- ΔR Uranium and thorium contents in different ceramic samples have been evaluated by using CR-39 and LR-115 type II solid state nuclear track detectors (SSNTDs) and calculating the probabilities for α -particles to reach and be registered on the SSNTD films. A new method has been developed based on calculating the stopping powers of the studied materials for the α - and β -particles emitted by the nuclei of the uranium and thorium series for evaluating annual $\alpha-$ and β dose rates in the considered ceramic samples. β -Dose rates due to potassium 40 (40K) have been evaluated for the studied materials. α -Dose rates of the considered ceramic samples have been compared with data obtained by using Bell's method.
- CC 71-7 (Nuclear Technology)

Section cross-reference(s): 57, 79

25656-90-0, Diethylene glycol bis(allyl carbonate) homopolymer ΙT RL: DEV (Device component use); USES (Uses)

> (CR-39; evaluating annual α - and β -dose rates emitted by U and Th series nuclei in different ceramic samples by using solid state nuclear track detectors)

ΙT 9004-70-0, LR 115II

RL: DEV (Device component use); USES (Uses)

(LR-115 type II; evaluating annual α - and β -dose rates emitted by U and Th series nuclei in different ceramic samples by using solid state nuclear track detectors)

7440-29-1, Thorium, analysis 14274-82-9, Thorium 228, analysis 14913-49-6, Bismuth 212, analysis 15389-34-1, Polonium 212, analysis

```
15756-58-8, Polonium 216, analysis 22481-48-7, Radon 220,
     analysis 28522-22-7, Radon 224, analysis
     RL: ANT (Analyte); ANST (Analytical study)
        (evaluating annual \alpha-dose rates emitted by Th
       series nuclei in different ceramic samples by using solid state nuclear
        track detectors)
                                  13966-29-5, Uranium 234, analysis
     7440-61-1, Uranium, analysis
     13981-52-7, Polonium 210, analysis 13982-63-3, Radium 226,
     analysis 14269-63-7, Thorium 230, analysis 14859-67-7, Radon
                   15422-74-9, Polonium 218, analysis 15735-67-8, Polonium
     222, analysis
     214, analysis
     RL: ANT (Analyte); ANST (Analytical study)
        (evaluating annual \alpha-dose rates emitted by U
        series nuclei in different ceramic samples by using solid state nuclear
        track detectors)
     14331-83-0, Actinium 228, analysis 14913-50-9, Thallium 208, analysis
ΙT
     15092-94-1, Lead 212, analysis 121239-98-3, Radon 228, analysis
     RL: ANT (Analyte); ANST (Analytical study)
        (evaluating annual \beta-dose rates emitted by Th series nuclei in
        different ceramic samples by using solid state nuclear track detectors)
     14255-04-0, Lead 210, analysis 14331-79-4, Bismuth 210, analysis
ΙΤ
     14733-03-0, Bismuth 214, analysis
                                       15035-09-3, Thallium 206, analysis
     15065-10-8, Thorium 234, analysis 15067-28-4, Lead 214, analysis
     15100-28-4, Protactinium 234, analysis 51634-37-8, Thallium 218,
     analysis
     RL: ANT (Analyte); ANST (Analytical study)
        (evaluating annual \beta-dose rates emitted by U series nuclei in
       different ceramic samples by using solid state nuclear track detectors)
                               THERE ARE 14 CITED REFERENCES AVAILABLE FOR THIS
REFERENCE COUNT:
                         14
                               RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
L84 ANSWER 10 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER:
                        1983:412475 CAPLUS Full-text
DOCUMENT NUMBER:
                        99:12475
ORIGINAL REFERENCE NO.: 99:1947a,1950a
TITLE:
                        Investigation of nitric acid for removal of noxious
                        radionuclides from uranium ore or mill tailings
                        Rvon, A. D.; Bond, W. D.; Hurst, F. J.; Scheitlin, F.
AUTHOR(S):
                        M.; Seeley, F. G.
                        Oak Ridge Natl. Lab., Oak Ridge, TN, 37830, USA
CORPORATE SOURCE:
SOURCE:
                        Uranium Mill Tailings Manage., Proc. Two NEA Workshops
                         (1982), Meeting Date 1981, 139-47. OECD: Paris, Fr.
                        CODEN: 49VJAB
DOCUMENT TYPE:
                        Conference
LANGUAGE:
                        English
     A conceptual process using HNO3 [7697-37-2] to extract 226Ra and 230Th in
     addition to the U from ore is proposed to decrease the potential hazard from
     discharge of mill tailings to the environment. This process removes ≤98% of
     the 226Ra and 230Th, yielding a residue containing as low as 10 pCi of Ra/q.
     Leaching of U from ores is consistently >99.5%. The residue after multistage
     leaching with HNO3 is resistant to further Ra leaching with water. Rn
     emanation from HNO3-leached residues generally is low due to the low Ra
     content. Heating to 800° causes further reduction of Rn emanation. Greater
     than 99% recovery of Ra from HNO3-leach solns. is obtained by carrying it on
     BaSO4. Good adsorption of Ra is also obtained on barite and Celite. Recovery
     of Th and U by solvent extraction using Bu3PO4 appears promising. Recycle of
     HNO3 may be accomplished by solvent extraction combined with evaporation and
     calcination.
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71-10 (Nuclear Technology)

CC

10/560922 Section cross-reference(s): 54 ΙT 10043-92-2, properties RL: PRP (Properties) (emanation of, from nitric acid leach solution of uranium ore) 7440-61-1P, properties 13981-52-7P, properties 13982-63-3P, ΤТ properties 14255-04-0P, properties 14269-63-7P, properties RL: RCT (Reactant); PREP (Preparation); RACT (Reactant or reagent) (leaching of, from uranium ore, in nitric acid solution) L84 ANSWER 11 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 1978:185371 CAPLUS Full-text DOCUMENT NUMBER: 88:185371 ORIGINAL REFERENCE NO.: 88:29107a,29110a Analysis of alpha emitters in the TITLE. coral, Favites virens, from Bikini Lagoon by solid-state track detection AUTHOR(S): Levy, Yitzhak; Miller, Donald S.; Friedman, Gerald M.; Noshkin, Victor E. Dep. Geol., Rensselaer Polytech. Inst., Troy, NY, USA CORPORATE SOURCE: Health Physics (1978), 34(3), 209-17 SOURCE: CODEN: HLTPAO; ISSN: 0017-9078 DOCUMENT TYPE: Journal LANGUAGE: English AB A quant. method for the nondestructive anal. of α -particle emitters in CaCO3 matrixes by solid-state track detection (cellulose nitrate) was developed. In an area of 4 mm2, .apprx.0.4 pCi/g can be measured routinely; smaller concns. can be determined but with a lower resolution Calibration methods used were (a) a Pu source of 0.15 μCi in conjunction with polycarbonate and CaCO3 absorbers of different thickness $(2-30 \mu m)$ and (b) a powdered coral sample from Enewetok Atoll that had been radiochem. analyzed for Pu radionuclides, 241Am, and other long-lived fission and activation products. Slabs of a coral, F. virens, from Bikini lagoon were analyzed. A quantity of the lphaparticle emitters detected in regions of the coral identified with growth during nuclear testing (1954, 1956, and 1959) are found in small discrete spots. Thin sections cut parallel to the direction of coral growth give different patterns of distributions. No such hot spots are evident in any post-test-yr growth sections although Pu and other long-lived fission and activation products were measured in these sections by radiochem. techniques. CC 8-1 (Radiation Biochemistry) alpha particle emitter detn coral ST ΙT Favites virens (alpha-particle emitters determination in, by solid-state track detector) 7440-29-1, analysis 7440-61-1, analysis 13981-16-3, analysis ΤТ 13981-52-7, analysis 13982-63-3, analysis 14119-33-6, analysis 14255-04-0, analysis 14269-63-7, analysis 14274-82-9, analysis 14596-10-2, analysis 15117-48-3, analysis 15117-96-1, analysis RL: ANT (Analyte); ANST (Analytical study) (determination of, by solid-state track detector) L84 ANSWER 12 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN

1977:570658 CAPLUS Full-text ACCESSION NUMBER: DOCUMENT NUMBER: 87:170658 ORIGINAL REFERENCE NO.: 87:26955a,26958a

TITLE: Emission and enrichments of radon daughters from Etna

volcano magma

Lambert, G.; Bristeau, P.; Polian, G. AUTHOR(S): Cent. Faibles Radioact., CNRS-CEA, Fr. CORPORATE SOURCE:

Geophysical Research Letters (1976), 3(12), 724-6 SOURCE:

CODEN: GPRLAJ; ISSN: 0094-8276

DOCUMENT TYPE: Journal LANGUAGE: English

AB Studies of the Etna volcano plume show that Rn daughters as well as gaseous Rn are directly emitted from the magma. Enrichment factors observed are 4 for Bi vs. Pb and 6 for Po vs. Pb. These enrichments correlate with relative volatility of the elements and/or their different compds. The 210Pb total output measured for the Etna plume is an insignificant source for this nuclide.

CC 53-3 (Mineralogical and Geological Chemistry)

IT 10043-92-2P, preparation 13981-52-7P, preparation

14255-04-0P, preparation RL: PREP (Preparation)

(emission of, from Etna Volcano)

L84 ANSWER 13 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 1958:14581 CAPLUS <u>Full-text</u>

DOCUMENT NUMBER: 52:14581
ORIGINAL REFERENCE NO.: 52:2594a-b

TITLE: Energy of α -particles from polonium-210

AUTHOR(S): Agapkin, I. I.; Gol'din, L. L.

SOURCE: Izvestiya Akademii Nauk SSSR, Seriya Fizicheskaya

(1957), 21, 909-12

CODEN: IANFAY; ISSN: 0367-6765

DOCUMENT TYPE: Journal LANGUAGE: Unavailable

The measurements were made with a magnetic α -spectrometer. Error sources were listed; it was observed that freshly made prepns. had reproducible results. By using a correction for the half-width of the line, obtained by varying the slit width, an energy of 5297.8 \pm 1.5 e.kv. was obtained, compared to the tabulated value of 5300.6 \pm 2.6 e.kv. For Em220 the energy was 6282.4 (tabulated value 6282.3 \pm 1.3 e.kv.).

CC 3A (Nuclear Phenomena)

IT 13981-52-7P, Polonium, isotope of mass 210

RL: PREP (Preparation)

(separation from Bi210 and Pb210)

IT 14255-04-0P, Lead, isotope of mass 210

RL: PREP (Preparation)

(separation from Bi210 and Po210)

IT 13981-52-7, Polonium, isotope of mass 210 22481-48-7, Radon,

isotope of mass 220 $(\alpha$ -rays from)

L84 ANSWER 14 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 1956:3794 CAPLUS Full-text

DOCUMENT NUMBER: 50:3794
ORIGINAL REFERENCE NO.: 50:690f-g

TITLE: Purification of radioactive deposits formed in radon

needles

AUTHOR(S): Ferreira de Miranda, C. CORPORATE SOURCE: Fac. cienc., Lisbon, Port.

SOURCE: Rev. fac. cienc. Univ. Lisboa (1954), 3, 191-5

DOCUMENT TYPE: Journal LANGUAGE: Unavailable

AB Rn needles are treated with 3N HNO3 and the liquid is centrifuged. The solution is heated up to 45° and a solution of 1.5N KCN is added. The precipitate is separated by centrifugation and redissolved in 0.5N HNO3. The procedure is repeated twice. The radioactive material formed by Ra D, Ra E, and Ra F is concentrated in the last HNO3 solution

CC 3A (Nuclear Phenomena)

IT 13981-52-7P, Polonium, isotope of mass 210 14255-04-0P,

Lead, isotope of mass 210
RL: PREP (Preparation)

(concentration from Rn deposits)

IT 10043-92-2, Radon (decay products of)

L84 ANSWER 15 OF 31 WPIX COPYRIGHT 2008 THOMSON REUTERS on STN

DUPLICATE 6

ACCESSION NUMBER: 1981-42085D [23] WPIX

TITLE: Locating deposits of uranium and thorium - by collecting

field samples forming chemical sons. and examining alpha

sensitive films exposed to delay prods.

DERWENT CLASS: K08; S03
INVENTOR: STIEFF L R

PATENT ASSIGNEE: (STIE-I) STIEFF L R

COUNTRY COUNT: 2

PATENT INFORMATION:

PATE	ENT NO	KIND	DATE	WEEK	LA	PG	MAIN	IPC
US 4	1268748	Α	19810519	(198123)*	ΕN			
CA 1	L145067	A	19830419	(198319)	ΕN			

APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE
US 4268748	A	US 1978-879578	19780221
US 4268748	A	US 1978-899937	19780425
US 4268748	A	US 1979-106424	19791221

INT. PATENT CLASSIF.:

IPC RECLASSIF.: G01V0005-00 [I,A]; G01V0005-00 [I,C]

ECLA: G01V0005-00 USCLASS NCLM: 250/255.000

NCLS: 250/472.100; 250/DIG.002

BASIC ABSTRACT:

US 4268748 A UPAB: 20050419 Uranium deposits are detected by collecting subsurface samples from a location and examining the samples for characteristic daughter prods. Pb, Bi and Po are chemically extracted from the samples. First and second alpha particle sensitive films are exposed to record decay of Po 214 and Po210. After etching, alpha particle damage populations are determined and correlated with sample locations in order to determine the position of a uranium deposit. Samples are collected from differentiated zones surrounding the deposit. Planchets on which alpha active Po from solution has been deposited are exposed to the alpha sensitive film. The method detects deposits of uranium by measuring Pb214, Bi 214, Po214, Pb210, and Po210 daughters. The deposits are distinguished from thorium deposits which can also be located. MANUAL CODE: CPI: K08-A; K09-J

EPI: S03-C03; S03-G01

L84 ANSWER 16 OF 31 WPIX COPYRIGHT 2008 THOMSON REUTERS on STN

ACCESSION NUMBER: 2003-776157 [73] WPIX CROSS REFERENCE: 2003-197783; 2003-898376

DOC. NO. CPI: C2003-213524 [73] DOC. NO. NON-CPI: N2003-621799 [73]

TITLE: Removal of radioactive contaminants from surface involves

retaining aqueous solution comprising wetting agent and active agent on surface and removing aqueous solution

containing radioactive contaminants from surface

DERWENT CLASS: A17; A25; A97; K07; P43

INVENTOR: MARTIN R T

PATENT ASSIGNEE: (BOBO-N) BOBOLINK INC

COUNTRY COUNT: 1

PATENT INFORMATION:

PATENT NO	KIND DATE	WEEK LA	PG	MAIN IPC
US 6605158	B1 20030812	(200373)* EN	1018	

APPLICATION DETAILS:

PATENT NO KIND	APPLICATION	DATE
US 6605158 B1 CIP of	US 2001-976467	20011012
US 6605158 B1	US 2002-283039	20021024

FILING DETAILS:

PATENT NO	KIND	PATENT NO
US 6605158 B1	CIP of	US 6497769 B

PRIORITY APPLN. INFO: US 2002-283039 20021024 US 2001-976467 20011012

INT. PATENT CLASSIF.:

IPC RECLASSIF.: C11D0011-00 [I,A]; C11D0011-00 [I,C]; C11D0007-02 [I,C];

C11D0007-08 [I,A]; C11D0007-22 [I,C]; C11D0007-26 [I,A]; C11D0007-32 [I,A]; G21F0009-28 [I,A]; G21F0009-28 [I,C]

ECLA: C11D0007-08; C11D0007-26E; C11D0007-32D; C11D0011-00B2D;

C11D0011-00B8; G21F0009-28

ICO: M11D0007:26A USCLASS NCLM: 134/010.000

NCLS: 134/002.000; 134/003.000; 134/022.190; 134/026.000; 134/028.000; 134/036.000; 134/041.000; 134/042.000; 210/682.000; 210/688.000; 376/308.000; 376/309.000; 376/310.000; 423/002.000; 423/003.000; 423/020.000; 510/110.000; 588/001.000; 588/020.000; 976/DIG.376;

976/DIG.391

BASIC ABSTRACT:

US 6605158 B1 UPAB: 20050601

NOVELTY - Radioactive contaminants are removed from a surface by applying to the surface an aqueous solution comprising a wetting agent and an active agent; retaining the aqueous solution on the surface for a time to allow radioactive contaminants to migrate into the solution; and removing the solution containing the radioactive contaminants from the surface. The active agent is a specified complex substituted keto-amine.

DETAILED DESCRIPTION - Removal of radioactive contaminants from a surface includes applying to the surface an aqueous solution comprising a wetting agent and an active agent; retaining the aqueous solution on the surface for a time to allow radioactive contaminants to migrate into the aqueous solution; and removing the aqueous solution containing the radioactive contaminants from the surface. The wetting agent is methanol, ethanol, propanol, isopropanol,

butanol, propargyl alcohol, tertiary butyl alcohol, pentanol, propylene glycol, polypropylene glycol, and/or ethylene glycol. The active agent is a complex substituted keto-amine having the formula (I). R-N(R') (CH2R'') (I)

R = abietyl, hydroabietyl, or dehydroabietyl; R'' = alpha ketonyl having fewer than 10C; and R' = H, or CH2R''.

USE - For removing radioactive contaminants from a surface.

ADVANTAGE - The inventive method has the effect of removing all of the radioactive contaminants from a previously contaminated surface. It is simple, economic, and an effective way of removing all of the radioactive contaminants from various surfaces or areas. MANUAL CODE:

CPI: A12-W11C; A12-W11F; K07-A03

ABEX EXAMPLE - Several decontamination solutions were prepared and tested utilizing various acids but containing the amount of keto-amine (0.5%), isopropyl alcohol (0.3%) and propargyl alcohol (0.3%). Each solution had a pH of at mostl.2. The solutions were applied as a low-pressure spray, allowed to remain at the site for 5 minutes and then removed by a squeegee. The spent solution was neutralized to a pH of 7, using a 50% sodium hydroxide solution. Surface contamination levels of both alpha and beta emissions were measured before and after treatment. The results showed degrees of effectiveness in removing radioactivity from surfaces utilizing a strong acid combined with a keto-amine as the active agent and a mixture of lower alcohols.

TECH

INORGANIC CHEMISTRY - Preferred Component: The aqueous solution further comprises an acid from hydrochloric acid, hydrofluoric acid, sulfuric acid, phosphoric acid, sulfurous acid, bromic acid, iodic acid, nitric acid, perchloric acid, oxalic acid, aqua regia, citric acid, sulfamic acid, glycolic acid, and/or ascorbic acid. It contains 0.1-1 wt.% wetting agent, 0.1-2 wt.% complex substituted keto-amine, and 0 wt.% acid. The complex substituted keto-amine has the molecular formula C33H45NO2C1H. The alpha ketonyl group is a ketone from acetone, methyl ethyl ketone, diacetone alcohol, isophorone, mesityl oxide, pentane dione, acetonyl acetone, cyclopentanone, cyclohexanone, or acetophenone. The wetting agent is a mixture of isopropanol and propargyl alcohol. INORGANIC CHEMISTRY - Preferred Component: The radioactive contaminant is a member of the lanthanide or actinide group. It is preferably Actinium-227, Americium-241, Americium-243, Antimony-124, Antimony-125, Barium-133, Beryllium-7, Bismuth-207, Cadmium-109, Calcium-45, Carbon-14, Cerium-139, Cerium-141, Cerium-144, Cesium-134, Cesium-135, Cesium-137, Chromium-51, Cobalt-56, Cobalt-57, Cobalt-58, Cobalt-60, Copper-67, Curium-242, Curium-243, Curium-244, Curium-247, Europium-152, Europium-154, Europium-155, Gadolinium-153, Germanium-68, Gold-195, Hafnium-181, Hydrogen-3 (Tritium), Iodine-125, Iodine-126, Iodine-129, Iodine-131, Iodine-133, Iridium-192, Iron-55, Iron-59, Lead-210, Manganese-54, Mercury-203, Neptunium-237, Nickel-59, Nickel-63, Niobium-94, Plutonium-236, Plutonium-238, Plutonium-239, Plutonium-240, Plutonium-241, Plutonium242, Plutonium-243, Plutonium-244, Polonium-210, Potassium-40, Promethium-147, Protactinium-231, Radium-223, Radium-224, Radium-226, Radium-228, Ruthenium-106, Samarium-151, Scandium-46, Selenium75, Silver-108m, Silver-110m, Sodium-22, Strontium-85, Strontium-89, Strontium-90, Sulfur-35, Tantalum-182, Technetium-99, Thallium-204, Thorium-natural, Thorium-228, Thorium-230, Thorium-232, Tin-113, Uranium-232, Uranium-233, Uranium-234, Uranium-235, Uranium-236, Uranium-238, Uranium-natural, Uranium-depleted, Yttrium-88, Yttrium-91, Zinc-65, Zirconium-95, or their associated decay products. INSTRUMENTATION AND TESTING - Preferred Method: The applying, retaining and removing steps are repeated to optimize decontamination. The aqueous solution is applied to the surface by spraying. It is retained on the surface for less than 10 minutes.

Preferred Component: The surface is a metal, plastic, glass, concrete, wood, fiberglass, fabric, and/or soil.

L84 ANSWER 17 OF 31 BIOSIS COPYRIGHT (c) 2008 The Thomson Corporation on

DUPLICATE 4

ACCESSION NUMBER: 1997:358484 BIOSIS Full-text

DOCUMENT NUMBER: PREV199799664887

TITLE: Retrospective assessment of radon exposure by

measurements of 210Po implanted in surfaces using an alpha

track detector technique.

Falk, R.; Mellander, H.; Nyblom, L.; Ostergren, I. AUTHOR(S):

Swedish Radiation Protection Inst., S-171 16 Stockholm, CORPORATE SOURCE:

Sweden

SOURCE: Environment International, (1996) Vol. 22, No. SUPPL. 1,

pp. S857-S861.

CODEN: ENVIDV. ISSN: 0160-4120.

DOCUMENT TYPE: Article LANGUAGE: English

Entered STN: 25 Aug 1997 ENTRY DATE:

Last Updated on STN: 25 Aug 1997

ABSTRACT: The radon exposure of the past is important in epidemiological studies where an assessment of lung cancer risk from indoor

exposure is evaluated. The long-lived decay product, ***radon***

210Pb (T/2 = 22 y), is implanted into indoor surfaces by alpha recoils

and can be monitored to give information about the previous radon

history. This gives an alternative or complementary method to the traditional measurements of the current average radon concentration.

Autoradiographic alpha-track methods to assess the 210Pb activity implanted in glass surfaces by measurement of 210Po alpha activity

were investigated to find a simple and reliable method for field use. One limiting factor at low exposure levels is the alpha background activity in the sub-surface material. In the search for a practical field method, the use of two different alpha-track detector materials was found successful. By exposing one Kodak LR-115 cellulose-nitrate film and one CR-39 detector side by side on glass panes, the background of the glass is measured with the LR-115 and both the background and the signal by the CR-39 detector. Results from

measurements in 31 dwellings show that an exposure of more than 1000 Bq cntdot y-m-3 to a glass surface can be measured with the (CR-LR) difference technique. Experiences from the field measurements show the method to be accurate, simple, and reliable and therefore a promising tool for future madom

epidemiological studies.

Radiation biology - Radiation effects and protective CONCEPT CODE:

measures 06506

Biochemistry studies - Minerals

Toxicology - Environment and industry 22506

Public health - Air, water and soil pollution 37015

Public health - Radiation health

INDEX TERMS: Major Concepts

> Biochemistry and Molecular Biophysics; Pollution Assessment Control and Management; Radiation Biology;

Toxicology

INDEX TERMS: Chemicals & Biochemicals

RADON; POLONIUM-210; CR-39

INDEX TERMS: Miscellaneous Descriptors

> ALPHA TRACK DETECTOR TECHNIQUE; ASSESSMENT METHOD; CARCINOGEN; CR-39 DETECTOR; EPIDEMIOLOGICAL STUDIES; EXPOSURE; INDOOR; LONG-LIVED DECAY PRODUCT; LUNG CANCER;

NEOPLASTIC DISEASE; POLLUTION; POLONIUM-210; RADIATION BIOLOGY; RADIONUCLIDES; RADON; RESPIRATORY SYSTEM DISEASE; RISK

REGISTRY NUMBER: 10043-92-2 (RADON)

13981-52-7 (POLONIUM-210)

25656-90-0Q (CR-39) 81283-55-8Q (CR-39)

L84 ANSWER 18 OF 31 BIOSIS COPYRIGHT (c) 2008 The Thomson Corporation on

TN DUPLICATE 5

ACCESSION NUMBER: 1989:159461 BIOSIS Full-text DOCUMENT NUMBER: PREV198987081562; BA87:81562

TITLE: POLONIUM IN FLORIDA USA GROUNDWATER AND ITS POSSIBLE

RELATIONSHIP TO THE SULFUR CYCLE AND BACTERIA.

AUTHOR(S): HARADA K [Reprint author]; BURNETT W C; LAROCK P A; COWART

JB

CORPORATE SOURCE: DEP OCEANOGRAPHY, FLA STATE UNIV, TALLAHASSEE, FLA 32306,

USA

SOURCE: Geochimica et Cosmochimica Acta, (1989) Vol. 53, No. 1, pp.

143-150.

CODEN: GCACAK. ISSN: 0016-7037.

DOCUMENT TYPE: Article FILE SEGMENT: BA LANGUAGE: ENGLISH

ENTRY DATE: Entered STN: 25 Mar 1989

Last Updated on STN: 25 Mar 1989

ABSTRACT: The last radioactive member of the 238U natural decay-series, ***210Po*** is normally considered a very particle-reactive isotope. Analysis of most natural waters shows that 210Po is present at very low activities, usually even lower than its insoluble precursor, 210Pb . We have recently discovered, however, that 210Pb exists at very high concentrations in groundwaters of some shallow aquifers in west central Florida. These waters tend to be fairly acidic (pH < 5), sulfide-bearing, and relatively high in 222Rn. Detailed study of one well with extraordinary levels 210Pb (.apprx. 1000 dpm/l) indicates that: (1) 210Po in this water is in great excess of radioactive equilibrium with its predecessors ***210Pb*** and 210Bi; (2) most Po in this water exists in a form which does not coprecipitate with an iron hydroxide scavenge; and (3) the conversion of soluble (0.2 μm filter) to particulate Po occurs over a time scale of a few days during sulfide oxidation. We suspect that Po cycling in this environment is related to the sulfur cycle and may, therefore, be

CONCEPT CODE: General biology - Conservation and resource management

00512

Radiation biology - General 06502

Circadian rhythms and other periodic cycles 07200

Ecology: environmental biology - Oceanography and limnology

07510

Biochemistry studies - Minerals 10069 Bacteriology, general and systematic 30000 Physiology and biochemistry of bacteria 31000

Food microbiology - Biodegradation and biodeterioration

39006

Soil science - Physics and chemistry 52805

INDEX TERMS: Major Concepts

influenced by sulfur bacteria.

Biochemistry and Molecular Biophysics;

Biosynchronization; Conservation; Estuarine Ecology (Ecology, Environmental Sciences); Radiation Biology;

Soil Science; Systematics and Taxonomy

INDEX TERMS: Miscellaneous Descriptors

AQUIFER CHEMICAL ANALYSIS RADON URANIUM

ORGANISM: Classifier

Bacteria 05000

Super Taxa

Microorganisms

Taxa Notes

Bacteria, Eubacteria, Microorganisms

REGISTRY NUMBER: 7440-08-6 (POLONIUM)

7704-34-9 (SULFUR) 10043-92-2 (RADON) 7440-61-1 (URANIUM)

L84 ANSWER 19 OF 31 BIOSIS COPYRIGHT (c) 2008 The Thomson Corporation on

STN

ACCESSION NUMBER: 2003:4191 BIOSIS <u>Full-text</u>

DOCUMENT NUMBER: PREV200300004191

TITLE: Experimental methods of determining the activity depth

distribution of implanted 210Pb in glass.

AUTHOR(S): Roos, Birgitta [Reprint Author]; Samuelsson, Christer

CORPORATE SOURCE: Department of Radiation Physics, Lund University Hospital,

SE-221 85, Lund, Sweden birgitta.roos@radfys.lu.se

SOURCE: Journal of Environmental Radioactivity, (2002) Vol. 63, No.

2, pp. 135-151. print.

ISSN: 0265-931X (ISSN print).

DOCUMENT TYPE: Article LANGUAGE: English

ENTRY DATE: Entered STN: 18 Dec 2002

Last Updated on STN: 18 Dec 2002

ABSTRACT: Glass is often used in radon surveys to estimate retrospective radon concentrations, as radon progenies are

embedded in the upper surface layer. Experimental methods based on etching to

determine the depth distribution of recoil-implanted 210Po in glass from radon decay in air is presented. By carefully controlling

chemical concentrations and exposure time during which the glass is etched, stepwise removal of the surface material was possible. Two different etching agents, diluted ${\it HF/HNO3}$ and NaOH were utilised, with very similar results.

Experimental recoil depths of 210Po agree with theoretical

calculations from the literature. The maximum implantation depth obtained using this procedure was $100+-20\,\mathrm{nm}$.

CONCEPT CODE: Radiation biology - General 06502

Biochemistry studies - General 10060

INDEX TERMS: Major Concepts

Biochemistry and Molecular Biophysics; Radiation Biology

INDEX TERMS: Chemicals & Biochemicals

glass; hydrofluoric acid; lead-210:

depth distribution; nitric acid; radon; sodium

hydroxide

INDEX TERMS: Methods & Equipment

radon survey: applied and field techniques

REGISTRY NUMBER: 7664-39-3 (hydrofluoric acid)

14255-04-0 (lead-210) 7697-37-2 (nitric acid) 10043-92-2 (radon)

1310-73-2 (sodium hydroxide)

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STN

ACCESSION NUMBER: 1985:424029 BIOSIS <u>Full-text</u> DOCUMENT NUMBER: PREV198580094021; BA80:94021

TITLE: A PRACTICAL METHOD FOR THE SIMULTANEOUS DETERMINATION OF

> THORIUM-234 RADON-226 LEAD-210 AND POLONIUM-210 IN SEAWATER.

AUTHOR(S): HARADA K [Reprint author]; TSUNOGAI S

DEP CHEM, FACULTY FISHERIES, HOKKAIDO UNIV, HAKODATE, 041, CORPORATE SOURCE:

JAPAN

SOURCE: Journal of the Oceanographical Society of Japan, (1985)

> Vol. 41, No. 2, pp. 98-104. CODEN: NKGKB4. ISSN: 0029-8131.

DOCUMENT TYPE: Article FILE SEGMENT: RΑ LANGUAGE: ENGLISH

ABSTRACT:A practical method was developed for the simultaneous determination of

226Ra, 234Th, 210Pb and 210Po in seawater. Samples are

spiked with 228Ra, 230Th, 208Po and common Pb to determine chemical yield.

These nuclides are coprecipitated with calcium carbonate and ferric from 20-50 l of seawater and separated by using ***hydroxide***

coprecipitation and ion exchange techniques. Counting sources of Ra and the other nuclides are prepared by electrodeposition onto silver discs. Their radioactivites are counted with an α -spectrometer and a low background

 β -counter. This method gives a standard deviation of .apprx. 5% for

replicate determination of 226Ra and other nuclides.

CONCEPT CODE: Methods - Field methods

> Mathematical biology and statistical methods 04500 Radiation biology - Radiation and isotope techniques

06504

Ecology: environmental biology - Oceanography

Biochemistry methods - General 10050 Biochemistry studies - General Biophysics - Methods and techniques

INDEX TERMS: Major Concepts

Biochemistry and Molecular Biophysics; Marine Ecology

(Ecology, Environmental Sciences); Methods and

Techniques; Radiology (Medical Sciences)

Miscellaneous Descriptors INDEX TERMS:

COPRECIPITATION ION EXCHANGE ELECTRODEPOSITION

RADIOACTIVITY SPECTROMETER

15065-10-8 (THORIUM-234) REGISTRY NUMBER:

> 16369-95-2 (RADON-226) 14255-04-0 (LEAD-210) 13981-52-7 (POLONIUM-210)

ANSWER 21 OF 31 ENERGY COPYRIGHT 2008 USDOE/IEA-ETDE on STN

ACCESSION NUMBER: 2008(2):11838 ENERGY Full-text

Uncertainties of retrospective radon concentration TITLE: measurements by multilayer surface trap detector.

Bastrikov, V.; Kruzhalov, A. (Ural State Technical AUTHOR:

Univ., Yekaterinburg (Russian Federation)); Zhukovsky,

M. (Institute of Industrial Ecology UB RAS,

Yekaterinburg (Russian Federation))

CORPORATE SOURCE: Societe Française de Radioprotection - SFRP, BP72,

92263 Fontenay-aux-Roses Cedex (France)

NUMBER OF REPORT: INIS-FR--7038

SOURCE: 2006. 11 p. Available from INIS in electronic form.

> Conference: Second European IRPA congress on radiation protection - Radiation protection: from knowledge to

action, Paris (France), 15-19 May 2006

DOCUMENT TYPE: Miscellaneous; Availability Note; Conference

COUNTRY: France LANGUAGE: English FIELD AVAILABILITY: AB

ABSTRACT: The detector for retrospective radon exposure measurements is developed. The detector consists of the multilayer package of solid-state nuclear track detectors LR-115 type. Nitrocellulose films works both as A-particle detector and as absorber decreasing the energy of A-particles. The uncertainties of implanted 210Pb measurements by two- and three-layer detectors are assessed in dependence on surface 210Po activity and gross background activity of the glass. The generalized compartment behavior model of radon decay products in the room atmosphere was developed and verified. It is shown that the most influencing parameters on the value of conversion coefficient from 210Po surface activity to average radon concentration are aerosol particles concentration, deposition velocity of unattached 218Po and air exchange rate. It is demonstrated that with the use of additional information on surface to volume room ratio, air exchange rate and aerosol particles concentration the systematic bias of conversion coefficient between surface activity of 210Po and average radon concentration can be decreased up to 30 %. (N.C.)

CLASSIFICATION CODE: *S46 INSTRUMENTATION RELATED TO NUCLEAR SCIENCE AND

TECHNOLOGY

NUMBER OF REPORT:

CONTROLLED TERM: ALPHA DECAY RADIOISOTOPES; LEAD 210; LEAD 214;

POLONIUM 214; POLONIUM 218; RADIATION DETECTORS; RADIATION DOSES; RADON; RISK ASSESSMENT; THORIUM 232;

RADIATION DOSES; RADON; RISK ASSESSMENT; THORTUM 23

URANIUM 238

BROADER TERM: ACTINIDE NUCLEI; ALPHA DECAY RADIOISOTOPES; BETA DECAY

RADIOISOTOPES; BETA-MINUS DECAY RADIOISOTOPES; DOSES;

ELEMENTS; EVEN-EVEN NUCLEI; FLUIDS; GASES; HEAVY NUCLEI; ISOTOPES; LEAD ISOTOPES; MEASURING

INSTRUMENTS; MICROSECONDS LIVING RADIOISOTOPES; MINUTES LIVING RADIOISOTOPES; NONMETALS; NUCLEI; POLONIUM ISOTOPES; RADIOISOTOPES; RARE GASES;

SPONTANEOUS FISSION RADIOISOTOPES; THORIUM ISOTOPES;

URANIUM ISOTOPES; YEARS LIVING RADIOISOTOPES

L84 ANSWER 22 OF 31 ENERGY COPYRIGHT 2008 USDOE/IEA-ETDE on STN

ACCESSION NUMBER: 2006(1):198 ENERGY Full-text

TITLE: 210Po and 210Pb in seals from the Baltic Sea and Lake

Saimaa, Finland.

AUTHOR: Solatie, D.; Rissanen, K. (Radiation and Nuclear

Safety Authority - STUK, Rovaniemi (Finland). Regional

Laboratory in Northern Finland); Vesterbacka, P. (Radiation and Nuclear Safety Authority - STUK, Helsinki (Finland). Natural Radiation Laboratory)

SSI--2005-15

SOURCE: Radiological Protection in Transition. Proceedings of

the 14. Regular Meeting of the Nordic Society for

Radiation Protection, NSFS.

Editor(s): Valentin, J.; Cederlund, T.; Drake, P.;
Finne, I.E.; Glansholm, A.; Jaworska, A.; Paile, W.;

Rahola, T.

Swedish Radiation Protection Authority, Stockholm

(Sweden)

Sep 2005. p. 237 of 386 p. Available from:

http://www.ssi.se/ssi/sub r/apporter/pdf/ssi/sub

r/app/sub 2/005/sub 1/5.pd; OSTI; Commercial

reproduction prohibited; OSTI as DE20674930; PURL: https://www.osti.gov/servlets/purl/20674930-Up6xtr/.

Conference: Radiological Protection in Transition. 14. Regular Meeting of the Nordic Society for Radiation Protection, NSFS, Raettvik (Sweden), 27 - 31 Aug 2005

ISSN: 0282-4434

DOCUMENT TYPE: Report Article; Conference

COUNTRY: Sweden LANGUAGE: English

FIELD AVAILABILITY: AB

ABSTRACT: 210Po and 210Pb are members of the 238U decay chain. 210Po is an alphaemitter with a half-life of 138 days, while its grandmother, 210Pb is a betaemitter with 22.3 year half-life. In the atmosphere 222Rn formats its decay products 210Po and 210Pb. These nuclides are deposited on to the surface of land and sea and thus enter the food chain. The naturally occurring radionuclides 210Po and 210Pb are important because their great contribution to radiation dose to human and other species. As top predators in the aquatic food chain, fish-eating seals are vulnerable to the accumulation of contaminants. In the Regional Laboratory in Northern Finland, measurements of 210Po and 210Pb activity concentrations in seals from the Baltic Sea and in ringed seals from Lake Saimaa have been performed. Concentrations of 210Po and 210Pb in seals were determined in muscle, liver, kidney and spleen. The results of 210Po and 210Pb activity concentrations and the ratio of 210Po / 210Pb in these samples are presented. (Summary-only contribution)

CLASSIFICATION CODE: *S54 Environmental sciences

CONTROLLED TERM: PINNIPEDS; BALTIC SEA; LAKES; POLONIUM 210; LEAD 210;

RADIOECOLOGICAL CONCENTRATION; MUSCLES; LIVER; SPLEEN;

KIDNEYS

BROADER TERM: ALPHA DECAY RADIOISOTOPES; ANIMALS; AQUATIC ORGANISMS;

BETA DECAY RADIOISOTOPES; BETA-MINUS DECAY RADIOISOTOPES; BODY; DAYS LIVING RADIOISOTOPES;

DIGESTIVE SYSTEM; ECOLOGICAL CONCENTRATION; EVEN-EVEN

NUCLEI; GLANDS; HEAVY NUCLEI; ISOMERIC TRANSITION

ISOTOPES; ISOTOPES; LEAD ISOTOPES; MAMMALS; NANOSECONDS LIVING RADIOISOTOPES; NUCLEI; ORGANS; POLONIUM ISOTOPES; RADIOISOTOPES; SEAS; SURFACE WATERS; VERTEBRATES; YEARS LIVING RADIOISOTOPES

ELEMENT TERM: Po; 210Po; is; Po is; Pb; 210Pb; Pb is; U; 238U; U is;

Rn; 222Rn; Rn is

L84 ANSWER 23 OF 31 ENERGY COPYRIGHT 2008 USDOE/IEA-ETDE on STN

ACCESSION NUMBER: 2004(14):81802 ENERGY Full-text

TITLE: Preparation of low level sealed 210Pb source for

random pulse generator.

AUTHOR: Mitsugashira, T.; Hara, M. (Tohoku Univ., Institute

for Materials Research, Oarai Branch, Oarai, Ibaraki

(Japan)); Tsuyuzaki, N. (OPTRANS Corp. (Japan))

NUMBER OF REPORT: KEK-PROC--2003-11

SOURCE: Proceedings of the fourth workshop on environmental

radioactivity.

Editor(s): Miura, Taichi

High Energy Accelerator Research Organization,

Tsukuba, Ibaraki (Japan)

Nov 2003. p. 263-268 of 384 p. 2 refs., 5 figs., 1 tab. Available from KEK(High Energy Accelerator Research Organization) 1-1 Oho, Tsukuba-shi,

Ibaraki-ken, 305-0801 JAPAN.

Conference: 4. workshop on environmental

radioactivity, Tsukuba, Ibaraki (Japan), 4 - 6 Mar

2003

DOCUMENT TYPE: Report Article; Conference; Availability Note

COUNTRY: Japan
LANGUAGE: Japanese

FIELD AVAILABILITY: AB

ABSTRACT: We have developed the random pulse generator (RPG) that utilizes alphaparticle detection with pin photodiode (PPD). In order to support an expected large market of RPG, a steady production system of weak (about 100 Bq) alpha sealed source is necessary, and, for such alpha-source, 210Pb-210Po source is the best

candidate on a viewpoint of environmental radioactivity impact. Two methods for such 210Pb-210Po sealed source preparation, namely direct deposition from isopropyl alcohol solution (IPA solution) of 210Pb-210Po nitrate (D-IPA method) and the 210Pb-210Po hydroxides precipitation (PPT method), were experimentally examined. In the former D-IPA method, an aliquot of IPA solution of 210Pb-210Po nitrate was directly dropped in a sealed cap for PPD and dried by heating. Then, a polycarbonate (PC) solution of 1/1 mixture of dichloromethane and dichloroethane was dropped on the source to make a thin (about 0.1 mg/cm2) film for radioactivity sealed. In the PPT method, 210Pb-210Po hydroxide was filtrated on a PC membrane filter (Nuclipore 0.1 mum) and the membrane filter was dissolved in a 1/1 mixture of dichloromethane and dichloroethane. The sealed 210Pb-210Po sources were prepared directly by dropping an aliquot of the solution into the PPD cap followed by its evaporation. The sealed sources thus prepared were subjected to 1m height fall-down test, air blowing test, and water immersion test. It was confirmed that no radioactive contaminants were coming off from the sealed source through these tests. (author)

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ELEMENT TERM:

CLASSIFICATION CODE: *S07 Isotopes and radiation sources

CONTROLLED TERM: ALCOHOLS; ALPHA DETECTION; COMPARATIVE EVALUATIONS;

DEPOSITION; FABRICATION; HYDROXIDES; LEAD 210; NITRATES; POLONIUM 210; PRECIPITATION; PULSE

GENERATORS; SEALED SOURCES

BROADER TERM: ALPHA DECAY RADIOISOTOPES; BETA DECAY RADIOISOTOPES;

BETA-MINUS DECAY RADIOISOTOPES; CHARGED PARTICLE DETECTION; DAYS LIVING RADIOISOTOPES; DETECTION; ELECTRONIC EQUIPMENT; EQUIPMENT; EVALUATION; EVEN-EVEN NUCLEI; FUNCTION GENERATORS; HEAVY NUCLEI; HYDROGEN COMPOUNDS; HYDROXY COMPOUNDS; ISOMERIC TRANSITION ISOTOPES; ISOTOPES; LEAD ISOTOPES; NANOSEC LIVING RADIOISOTOPES; NITROGEN COMPOUNDS; NUCLEI; ORGANIC COMPOUNDS; OXYGEN COMPOUNDS; POLONIUM ISOTOPES;

RADIATION DETECTION; RADIATION SOURCES; RADIOISOTOPES;

SEPARATION PROCESSES; YEARS LIVING RADIOISOTOPES Pb; 210Pb; is; Pb is; Pb*Po; Pb sy 2; sy 2; Po sy 2;

Po is; 210Po; 210Pb-210Po; D

L84 ANSWER 24 OF 31 ENERGY COPYRIGHT 2008 USDOE/IEA-ETDE on STN

ACCESSION NUMBER: 1994(14):96992 ENERGY Full-text

TITLE: Evaluation of radionuclide levels and radiological

dose in three populations of marine mammals in the $% \left(1\right) =\left(1\right) \left(1\right)$

eastern Canadian Arctic.

AUTHOR: Macdonald, C.R.; Ewing, L.L.; Wiewel, A.M.; Harris,

D.A. (AECL Research, Pinawa, Manitoba (Canada)); Stewart, R.E.A. (Dept. of Fisheries and Oceans, Winnipeg, Manitoba (Canada). Freshwater Inst.)

NUMBER OF REPORT: CONF-931152--

SOURCE: Ecological risk assessment: Lessons learned?. Abstract

book. Anon.

Pensacola, FL: Society of Environmental Toxicology and

Chemistry. 1993. p. 254 of 356 p. Society of

Chemistry, 1995. p. 254 of 556 p. Society of

Environmental Toxicology and Chemistry Office, 1010 North 12th Avenue, Pensacola, FL 32501-3307 (United

States).

Conference: 14. annual meeting of the Society of Environmental Toxicology and Chemistry (SETAC), Houston, TX (United States), 14-18 Nov 1993

DOCUMENT TYPE: Book Article; Conference

COUNTRY: United States

LANGUAGE: English
FIELD AVAILABILITY: AB

ABSTRACT: Radionuclide levels were measured in beluga, walrus and ringed seal populations collected in 1992 to assess radiation dose and changes in dose with age and sex. The authors hypothesized that Arctic marine food chains accumulate high levels of naturally-occurring radionuclides such as polonium-210 and that radiation may pose a stress to animals which also accumulate metals such as cadmium. Liver, kidney, muscle and jawbone were analyzed by gamma spectrometry for cesium-137, cesium-134, lead-210, potassium-40 and radium-226 and fission-derived nuclides. Polonium-210 was analyzed by alpha spec after autodeposition onto a silver disk. Cesium-137 concentrations in muscle in all three populations were low, and ranged from below detection limits to 10 Bg/kg ww. There was no evidence of fissionderived radionuclides such as zinc-65 or cobalt-60. Lead-210 levels ranged from below detection limits in muscle of ringed seal and walrus to a mean of 82.3 Bq/kq ww in walrus bone. Polonium-210 in the three population ranged from 10 to 30 Bq/kg ww in bone and kidney. The major contributor to dose in the animals was polonium-210 because it is an alpha emitter and accumulates to moderate levels in liver and kidney. Radiological dose is approximately 20--30 times higher than background in humans, and is considerably lower than the dose observed in terrestrial food chains *560162; C2120 in the Arctic CLASSIFICATION CODE:

CONTROLLED TERM:

AQUATIC ORGANISMS; CESIUM 134; CESIUM 137; CONTAMINATION; FOOD CHAINS; LEAD 210; MAMMALS; NATURAL

RADIOACTIVITY; POLONIUM 210; POTASSIUM 40; RADIUM 226; SENSITIVITY; TISSUE DISTRIBUTION

*MAMMALS: *SENSITIVITY; *FOOD CHAINS: *CONTAMINATION;

*CESIUM 137: *TISSUE DISTRIBUTION; *CESIUM 134:

*TISSUE DISTRIBUTION; *LEAD 210: *TISSUE DISTRIBUTION;

*POTASSIUM 40: *TISSUE DISTRIBUTION; *RADIUM 226:

*TISSUE DISTRIBUTION; *POLONIUM 210: *TISSUE

DISTRIBUTION

BROADER TERM:

ALKA; ALKALI METAL ISOTOPES; ALKALINE EARTH ISOTOPES; ALPHA DECAY RADIOISOTOPES; ANIMALS; BETA DECAY RADIOISOTOPES; BETA-MINUS DECAY RADIOISOTOPES; BETA-PLUS DECAY RADIOISOTOPES; CARBON 14 DECAY RADIOISOTOPES; CESIUM ISOTOPES; DAYS LIVING RADIOISOTOPES; DISTRIBUTION; ELECTRON CAPTURE RADIOISOTOPES; EVEN-EVEN NUCLEI; HEAVY ION DECAY RADIOISOTOPES; HEAVY NUCLEI; HOURS LIVING

RADIOISOTOPES; INTERMEDIATE MASS NUCLEI; INTERNAL CONVERSION RADIOISOTOPES; ISOMERIC TRANSI; ISOMERIC TRANSITION ISOTOPES; ISOTOPES; LEAD ISOTOPES; LIGHT NUCLEI; NANOSEC LIVING RADIOISOTOPES; NUCLEI; ODD-EVEN NUCLEI; ODD-ODD NUCLEI; POLONIUM ISOTOPES; POTASSIUM

ISOTOPES; RADIOACTIVITY; RADIOISOTOPES; RADIUM ISOTOPES; VERTEBRATES; YEARS LIVING RADIOISOTOPES

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ACCESSION NUMBER: 1985(18):134853 ENERGY Full-text

TITLE: Application of dual NaI-CsI(T1) detectors to the in

vivo detection and localization of radon seeds.

AUTHOR: Anon. [United States]

CORPORATE SOURCE: New York Univ., NY (USA). Inst. of Environmental

Medicine (4664000)

NUMBER OF REPORT: DOE/EV/04326--6; DE85001413

SOURCE: In vivo measurements of bone-seeking radionuclides.

Progress report, 1981-1984.

Cohen, N.

1984. pp. 89-96 Availability: NTIS, PC A06/MF A01; 1.

DOCUMENT TYPE: Report Article COUNTRY: United States

LANGUAGE: English

DOCUMENT NUMBER: ERA-10:043005

ABSTRACT: Sealed gold capillary tubes containing radon have been used as implants for the treatment of certain tumors and lesions. The short-lived daughters of Rn-222 decay with the 3.82 day half-life of the parent. The seeds are generally left after treatment since it is believed that the residual activity from Pb-210, Bi-210 and Po-210 does not represent a significant radiological hazard to the patient. Some have suggested that chronic low-level irradiation of the implantation site may result in the formation of a tumor. Radon seeds which were implanted more than 20 years ago are now being considered for removal. Three subjects were measured in our whole body counting facility using NaI-CsI(T1) detectors. With the NaI-CsI(T1) detector placed directly over the implantation area, it was possible to observe the K X-rays characteristic of gold which are produced as a result of beta particle interaction with the gold casing. 2 refs., 3 figs., 1 tab. CLASSIFICATION CODE: *560151; C1500

CONTROLLED TERM:

*RADIATION SOURCE IMPLANTS: *RADIATION HAZARDS; *RADON 222: *RADIATION SOURCE IMPLANTS; BISMUTH 210; BREMSSTRAHLUNG; DAUGHTER PRODUCTS; GOLD; JAW; LEAD 210; LEGS; MAMMARY GLANDS; NEOPLASMS; RADIOTHERAPY; SOLID SCINTILLATION DETECTORS; WHOLE-BODY COUNTERS; X RADIATION

BROADER TERM:

ALPHA DECAY RADIOISOTOPES; BETA DECAY RADIOISOTOPES;
BETA-MINUS DECAY RADIOISOTOPES; BISMUTH ISOTOPES;
BODY; BODY AREAS; DAYS LIVING RADIOISOTOPES; DISEASES;
ELECTROMAGNETIC RADIATION; ELEMENTS; EVEN-EVEN NUCLEI;
GLANDS; HAZARDS; HEALTH HAZARDS; HEAVY NUCLEI;
IMPLANTS; IONIZING RADIATIONS; ISOTOPES; LEAD
ISOTOPES; LIMBS; MEASURING INSTRUMENTS; MEDICINE;
METALS; NUCLEAR MEDICINE; NUCLEI; ODD-ODD NUCLEI;
ORGANS; RADIATION DETECTORS; RADIATION SOURCES;
RADIATIONS; RADIOISOTOPES; RADIOLOGY; RADON ISOTOPES;
SCINTILLATION COUNTERS; SKELETON; SKULL; THERAPY;
TRANSITION ELEMENTS; YEARS LIVING RADIOISOTOPES
CS*I*Na*T; CS sy 4; sy 4; I sy 4; Na sy 4; T sy 4;
NaI; Na cp; cp; I cp; CsI(T; Cs cp; T cp; NaI-CsI(T; Rn; Pb; Bi; Po; K

ELEMENT TERM:

L84 ANSWER 26 OF 31 ENERGY COPYRIGHT 2008 USDOE/IEA-ETDE on STN

ACCESSION NUMBER: 1983(10):83361 ENERGY Full-text

TITLE: Field method for detecting deposits containing uranium

and thorium. (Patent.)

AUTHOR: Stieff, L.R. [United States]

PATENT INFORMATION: US 4336451 22 Jun 1982

v p. PAT-APPL-106424.

APPLICATION INFORMATION: 21 Dec 1979

DOCUMENT TYPE: Patent

COUNTRY: United States

LANGUAGE: English

ABSTRACT: Locations of buried deposits are determined by detecting

the presence of 2exclamation4PB, 214BI, 214PO, 210BP, 210BI and 210PO in solutions obtained by chemically leaching these elements from rocks and soil. Polonium from the solution is plated on silver foil planchets. Alpha sensitive films are exposed to the alpha decay of 214PO and 210PO by contacting the films with the planchets. The films, when etched, reveal the damage caused by the passage of the high energy alpha particles. Alpha damage as a function of sample size, volume of solution used, planchet, foil or film area and exposure times measures concentration of 214PO and 210PO in the sample. Anomalus concentrations suggest presence of buried deposits containing uranium. Similar anomalus

concentrations of alpha damage from 212BI and 212PO formed in films exposed to foils or planchets plated from leached solutions containing 212PB, 212BI and 212PO suggest deposits containing thorium. Plotting normalized alpha damage population (Numbers of alpha per gram, per square mm., per hour) and sample locations suggest mineral deposit locations.

INT. PATENT CLASSIF.: G01V005-00

CLASSIFICATION CODE: *050200; 050100; B3100

CONTROLLED TERM: *URANIUM DEPOSITS: *GEOPHYSICAL SURVEYS; ALPHA

DETECTION; BISMUTH 210; BISMUTH 214; LEACHATES; LEAD 210; LEAD 214; POLONIUM 210; POLONIUM 214; QUANTITY

RATIO: THORIUM

BROADER TERM: ACTINIDES; ALPHA DECAY RADIOISOTOPES; BETA DECAY

RADIOISOTOPES; BETA-MINUS DECAY RADIOISOTOPES; BISMUTH

ISOTOPES; CHARGED PARTICLE DETECTION; DAYS LIVING RADIOISOTOPES; DETECTION; DISPERSIONS; ELEMENTS; EVEN-EVEN NUCLEI; GEOLOGIC DEPOSITS; HEAVY NUCLEI; ISOTOPES; LEAD ISOTOPES; METALS; MICROSEC LIVING RADIOISOTOPES; MINERAL RESOURCES; MINUTES LIVING RADIOISOTOPES; MIXTURES; NUCLEI; ODD-ODD NUCLEI;

POLONIUM ISOTOPES; RADIATION DETECTION; RADIOISOTOPES; RESOURCES; SECONDS LIVING RADIOISOTOPES; SOLUTIONS;

SURVEYS; YEARS LIVING RADIOISOTOPES

ELEMENT TERM: B*I; BI; 214BI; is; B is; 214B; B cp; cp; I cp; O*P;

PO; 214PO; P is; 214P; P cp; O cp; B*P; BP; 210BP; 210B; 210BI; 210PO; 210P; 212BI; 212B; 212PO; 212P

L84 ANSWER 27 OF 31 ENERGY COPYRIGHT 2008 USDOE/IEA-ETDE on STN

ACCESSION NUMBER: 1982(11):85211 ENERGY

TITLE: Airborne radiological sampling of Mount St. Helens

plumes.

AUTHOR: Andrews, V.E. [United States]

CORPORATE SOURCE: Office of Radiation Programs, Las Vegas, NV (USA)

(9511167)

NUMBER OF REPORT: PB--81-213795

Apr 1981. 20 p. Availability: NTIS, PC A02/MF A01.

DOCUMENT TYPE: Report

COUNTRY: United States

LANGUAGE: English

DOCUMENT NUMBER: ERA-07:034689

ABSTRACT: Particulate and gaseous samples for radiologial analyses were collected from the plumes created by eruptions of Mount St. Helens. The sampling aircraft and equipment used are routinely employed in aerial radiological surveillance at the Nevada Test Site by the Environmental Protection Agency's Environmental Monitoring Systems Laboratory in Las Vegas, Nevada. An initial sample set was collected on April 4, 1980, during the period of recurring minor eruptions. Samples were collected again on May 19 and 20 following the major eruption of May 18. The Environmental Protection Agency's Office of Radiation Programs analyzed the samples for uranium and thorium isotopes, radium-226, lead-210, polonium-210, and radon-222. Other laboratories analyzed samples to determine particle size distribution and elemental composition. The only samples containing radioactivity above normal ambient levels were collected on May 20. Polonium-210 concentrations in the plume, determined from a sample collected between 5 and 30 km from the crater, were approximately an order of magnitude above background. Radon-222 concentrations in samples collected from the plume centerline at a distance of 15 km averaged approximately four times the average surface concentrations. The small increases in radioactivity would cause no observable adverse health effects.

CLASSIFICATION CODE: *500300; 510301

CONTROLLED TERM: *LAVA: *CHEMICAL COMPOSITION; *VOLCANIC GASES:

*CHEMICAL COMPOSITION; *MT ST HELENS: *LAVA; *MT ST HELENS: *RADIATION MONITORING; *LAVA: *RADIATION MONITORING; *VOLCANIC GASES: *RADIATION MONITORING; *MT ST HELENS: *VOLCANIC GASES; AERIAL SURVEYING; AIR POLLUTION; DISTRIBUTION; LEAD; LEAD 210; PARTICLE SIZE; PLUMES; POLONIUM; POLONIUM 210; RADIOACTIVE AEROSOLS; RADIOACTIVE MATERIALS; RADIOACTIVITY; RADIUM; RADIUM 226; RADON; RADON 222; SAMPLING; THORIUM ISOTOPES; URANIUM ISOTOPES

BROADER TERM:

ACTINIDE ISOTOPES; AEROSOLS; ALKALINE EARTH ISOTOPES; ALKALINE EARTH METALS; ALPHA DECAY RADIOISOTOPES; BETA DECAY RADIOISOTOPES; BETA-MINUS DECAY RADIOISOTOPES; COLLOIDS; DAYS LIVING RADIOISOTOPES; DISPERSIONS; ELEMENTS; EVEN-EVEN NUCLEI; FLUIDS; GASES; HEAVY NUCLEI; ISOTOPES; LEAD ISOTOPES; MATERIALS; METALS; MONITORING; MOUNTAINS; NONMETALS; NUCLEI; POLLUTION; POLONIUM ISOTOPES; RADIOISOTOPES; RADIUM ISOTOPES; RADIOISOTOPES; RADIOISOTOPES; RADIOISOTOPES

L84 ANSWER 28 OF 31 ENERGY COPYRIGHT 2008 USDOE/IEA-ETDE on STN

ACCESSION NUMBER: 1983(1):7074 ENERGY Full-text

TITLE: Focusing chromatography using a dilute precipitation

reagent. Separation of 226Ra decayed nuclides by

potassium fluoride.

AUTHOR: Furushima, K.; Shinaqawa, M. (Kinki Univ.,

Higashi-Osaka, Osaka (Japan). Faculty of Science and

Technology) [Japan]

SOURCE: Radioisotopes (Tokyo) (Jun 1981) v. 30(6) p. 299-304

CODEN: RAISAB ISSN: 0033-8303

DOCUMENT TYPE: Journal COUNTRY: Japan LANGUAGE: Japanese

ABSTRACT: The electrophoresis of 226Ra and it's decayed nuclides was carried out by using hydrochloric acid in the positive electrode cell and aqueous solution of potassium fluoride in the negative electrode cell. Experimental factors, i.e. concentrations of potassium fluoride and hydrochloric acid, intensity of electric field, duration for electrophoresis and pH value of the potassium fluoride solution etc. were examined. The sample solution was of 0.01 M hydrochloric acid with a tracer amount of 226Ra and 210Pb. Each of these nuclides was carrier free and was 4GBq/l(lmuCi/ml) in concentration. For the sake of autoradiography, a photoengraving film (Fujilith Ortho Film, TAC = 135) was made use of examining the locations of the separated radioactive bands obtained on the paper strip. Their beta and gamma radioactivities were detected as the film darkening on developing the film. According to the positions thus detected, the strip filter paper was cut into pieces and the species of radionuclides were determined by the measurements of energies and half-lives on beta and gamma rays. The locations of the separated alpha-radionuclides were decided by etching the surface of it's film with 6 M aqueous solution of sodium hydroxide and the species of radionuclides were determined by the solid state alpha-track detection method. As the result, the optimum conditions for the separation are to use 0.01 M hydrochloric acid in the positive CLASSIFICATION CODE: *400703; B1300

CONTROLLED TERM: *RADIUM 226: *CHROMATOG

*RADIUM 226: *CHROMATOGRAPHY; AUTORADIOGRAPHY; DAUGHTER PRODUCTS; HYDROCHLORIC ACID; LEAD 210; PH VALUE; POTASSIUM FLUORIDES; PRECIPITATION; QUANTITY RATIO; ELECTROPHORESIS; AQUEOUS SOLUTIONS; ELECTRODES;

BISMUTH 210; POLONIUM 210; LEAD 214
Ra: 226Ra: is: Ra is: Ph: 210Ph: Ph is

ELEMENT TERM: Ra; 226Ra; is; Ra is; Pb; 210Pb; Pb is

L84 ANSWER 29 OF 31 EMBASE COPYRIGHT (c) 2008 Elsevier B.V. All rights

reserved on STN

AUTHOR:

ACCESSION NUMBER: 1997217207 EMBASE Full-text

TITLE: Retrospective assessment of radon exposure by

measurements of (210)Po implanted in

surfaces using an alpha track detector technique. Falk, R. (correspondence); Mellander, H.; Nyblom, L.;

Ostergren, I.

CORPORATE SOURCE: Swed. Radiation Protection Institute, S-171 16 Stockholm,

Sweden.

AUTHOR: Falk, R. (correspondence)

CORPORATE SOURCE: Swedish Radiation Protection Inst., S-171 16 Stockholm,

Environment International, (1997) Vol. 22, No. SUPPL. 1, SOURCE:

pp. S857-S861.

Refs: 4

ISSN: 0160-4120 CODEN: ENVIDV

PUBLISHER IDENT.: S 0160-4120(96)00193-6

United Kingdom COUNTRY:

DOCUMENT TYPE: Journal; Conference Article; (Conference paper) FILE SEGMENT: 046 Environmental Health and Pollution Control

LANGUAGE: English English SUMMARY LANGUAGE:

ENTRY DATE: Entered STN: 22 Aug 1997

Last Updated on STN: 22 Aug 1997

The radon exposure of the past is important in epidemiological studies ABSTRACT: where an assessment of lung cancer risk from indoor

radon exposure is evaluated. The long-lived decay product, (***210***)Pb (T 1/4 = 22 y), is implanted into indoor surfaces by

alpha recoils and can be monitored to give information about the previous ***radon*** history. This gives an alternative or complementary method to

the traditional measurements of the current average radon

concentration. Autoradiographic alpha-track methods to assess the (210

)Pb activity implanted in glass surfaces by measurement of (

210)Po alpha activity were investigated to find a simple and reliable method for field use. One limiting factor at low exposure levels is the alpha background activity in the sub-surface material. In the search for a practical field method, the use of two different alpha-track detector materials was found successful. By exposing one Kodak LR-115 cellulose-nitrate ***film*** and one CR- 39 detector side by side on glass panes, the background of the glass is measured with the LR-115 and both the background and

the signal by the CR-39 detector. Results from measurements in 31 dwellings show that an exposure of more than 1000 Bq.ovrhdot.y.ovrhdot.m(-3) to a glass surface can be measured with the (CR- LR) difference technique. Experiences from the field measurements show the method to be accurate, simple, and reliable and therefore a promising tool for future radon

CONTROLLED TERM: Medical Descriptors:

epidemiological studies.

alpha radiation autoradiography *cancer risk conference paper

*environmental exposure

priority journal reliability

retrospective study

CONTROLLED TERM: Drug Descriptors:

*radon

CAS REGISTRY NO.: (radon) 10043-92-2

L84 ANSWER 30 OF 31 COMPENDEX COPYRIGHT 2008 EEI on STN ACCESSION NUMBER: 1984(11):201224 COMPENDEX Full-text

DOCUMENT NUMBER: 8411121412

; *8491369

TITLE: STUDY OF 210Pb AND 210Po

DISTRIBUTIONS IN ENVIRONMENTAL SAMPLES BY CR-39 TRACK

DETECTOR.

AUTHOR: Hunyadi, I. (Hungarian Acad of Sciences, Inst of

Nuclear Research, Debrecen, Hung); Somogyi, G.;

Szilagyi, S.

SOURCE: Nucl Tracks Radiat Meas v 8 n 1-4 1984, Solid State

Nucl Track Detect, Proc of the Int Conf, 12th,

Acapulco, Mex, Sep 4-10 1983 p 491-495

SOURCE: Nucl Tracks Radiat Meas v 8 n 1-4 1984, Solid State

Nucl Track Detect, Proc of the Int Conf, 12th,

Acapulco, Mex, Sep 4-10 1983 p 491-495

CODEN: NUTRDQ ISSN: 0191-278X

PUBLICATION YEAR: 1983 LANGUAGE: English

ABSTRACT: Activity concentration distributions of long-lived alpha-emitters in aerosol samples are analyzed by high-resolution autoradiography in CR-39.A study of the alpha-activity attached to aerosols of different particulate sizes separated by a cascade impactor is also performed. It is found that, in the majority of samples, the alpha-activity can be dominantly related to the presence of Po-210 produced by its beta-active precursor Pb-210. Analysis of alpha-decay properties was done by autoradiographs taken at different post-sampling times. Spectroscopic studies of individual alpha tracks and track clusters were developed for high resolution alpha energy determination. Measured parameters were the major axis of surface track opening, the diameter of etched out track end, the total length measurable on the surface along the projected track, and the thickness of the layer etched away from the detector surface. 5 refs. CLASSIFICATION CODE: 944 Moisture, Pressure & Temperature, & Radiation

Measuring Instruments; 443 Meteorology; 622 Radioactive Materials; 815 Plastics & Polymeric Materials; 932 High Energy, Nuclear & Plasma Physics

CONTROLLED TERM: *PARTICLE DETECTORS:Applications; ATMOSPHERIC

RADIOACTIVITY: Analysis; POLYCARBONATES

SUPPLEMENTARY TERM: CR-39; AUTORADIOGRAPHY; ENVIRONMENTAL ALPHA ACTIVITY;

SSNTD; SOLID STATE NUCLEAR TRACK DETECTORS

ELEMENT TERM: Po; Pb; 210Pb; is; Pb is; 210Po; Po is

L84 ANSWER 31 OF 31 SCISEARCH COPYRIGHT(c) 2008 The Thomson Corporation on STN

ACCESSION NUMBER: 1994:283522 SCISEARCH Full-text

THE GENUINE ARTICLE: NJ812

TITLE: AN EFFICIENT QUANTITATIVE TECHNIQUE FOR THE SIMULTANEOUS

ANALYSES OF RADON DAUGHTERS PB-

210, BI-210 AND PO-210

AUTHOR: CHURCH T M (Reprint); HUSSAIN N; FERDELMAN T G; FOWLER S W

CORPORATE SOURCE: UNIV DELAWARE, COLL MARINE STUDIES, NEWARK, DE 19716

(Reprint); IAEA, ENVIRONM LAB, MONACO 98012, MONACO

COUNTRY OF AUTHOR: USA; MONACO

SOURCE: TALANTA, (FEB 1994) Vol. 41, No. 2, pp. 243-249.

ISSN: 0039-9140.

PUBLISHER: ELSEVIER SCIENCE BV, PO BOX 211, 1000 AE AMSTERDAM,

NETHERLANDS.

DOCUMENT TYPE: Article; Journal

FILE SEGMENT: PHYS
LANGUAGE: English
REFERENCE COUNT: 29

ENTRY DATE: Entered STN: 1994

Last Updated on STN: 1994

ABSTRACT:

An improved and time efficient technique has been developed for quantitative determination of the long-lived Rn-222 daughters (Pb-210, ***Po*** -210 and Bi-209) in atmospheric and oceanic samples. sample is first spiked with yield tracers for polonium (208 or 209), bismuth (207), and lead (stable lead carrier). These nuclides may then be scavenged through iron hydroxide precipitation and redissolved in a dilute (pH approximately 2) nitric acid plating medium with citrate and hydroxylamine hydrochloride at 90-degrees centrigrade with constant stirring. silver planchet is suspended in the solution which plates polonium to high Second, a nickel planchet is suspended in the same solution which efficiency. is maintained hermetic (e.g. bubbling with helium) and bismuth is plated next with high efficiency. Third, lead is purified from the same solution using anion exchange techniques and isolated for beta counting as the sulfate. Polonium is analyzed by isotope dilution alpha spectrometry. Bismuth and lead are analyzed by anti-coincident beta counting in a low level shield. case of bismuth, the 207 tracer is added in quantities at least comparable to the background of the beta system such that counting before and after the decay of 2 Bi-210 gives the bismuth yield.

The unique characteristics of this technique are its speed and efficiency; all three radon daughters can be isolated for counting within 4 hr of pre-treating the sample. The remaining solution can be treated subsequently for other analyses as appropriate.

CATEGORY: CHEMISTRY, ANALYTICAL

SUPPL. TERM PLUS: GROUNDWATERS; SEPARATION; POLONIUM

REFERENCE(S):

Referenced Author (RAU)	(RPY) (RV	L) (RPG)	(RWK)
ANON		·	IEXPT NUCLEAR SCI
· -			GEOCHIM COSMOCHIM AC
BAGNALL, K W	11957		CHEM RARE RADIOELEME
			HLTH PHYS
•	11966 38	1189	ANAL CHEM
•		•	CR HEBD ACAD SCI
·			GEOCHIM COSMOCHIM AC
	1974 5		
	1989 53	1143	GEOCHIM COSMOCHIM AC
HELKAMP, R W	1979 30	237	INT J APPL RADIAT IS
HELKAMP, R W HUSSAIN, N HUSSAIN, N HUSSAIN, N	1982 58	430	EARTH PLANET SC LETT
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POET, S E	1972 77	6515	J GEOPHYS RES
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TUREKIAN, K K	1977 5	1227	ANNU REV EARTH PL SC
VONOETTINGEN, W F	1930 10	221	PHYSIOL REV
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L3
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L4
          4376 SEA ABB=ON L3
L5
           1219 SEA ABB=ON L4 AND L5
1.6
             28 SEA ABB=ON L2/P AND L3/P
L7
                E RADIATION SOURCES+ALL/CT
L8
           2364 SEA ABB=ON RADIATION SOURCES/CT
L9
              4 SEA ABB=ON L6 AND L8
               D SCAN TI
L10
        903425 SEA ABB=ON A/OBI
L11
           118 SEA ABB=ON L10 AND L6
L12
          6038 SEA ABB=ON L10(L)(SOURCE#/OBI OR EMIT?/OBI)
            28 SEA ABB=ON L12 AND L6
L13
             1 SEA ABB=ON L12 AND L7
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        258343 SEA ABB=ON SEAL?/BI
L15
L16
              2 SEA ABB=ON (L13 OR L7) AND L15
                D SCAN TI
        881850 SEA ABB=ON FILM#/OBI
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L18
         76855 SEA ABB=ON POLYCARBONATE#/BI
        353488 SEA ABB=ON HYDROXIDE#/BI
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L22
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L23
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L24
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L26
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L27
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L29
L30
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             5 SEA ABB=ON L5(L)PUR/RL
L34
             2 SEA ABB=ON L33 AND L34
L35
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INDEX '1MOBILITY, 2MOBILITY, ABI-INFORM, ADISCTI, AEROSPACE, AGRICOLA, ALUMINIUM, ANABSTR, ANTE, APOLLIT, AQUALINE, AQUASCI, AQUIRE, BABS, BIBLIODATA, BIOENG, BIOSIS, BIOTECHABS, BIOTECHDS, BIOTECHNO, CABA, CAOLD, CAPLUS, CASREACT, CBNB, CEABA-VTB, CERAB, ...' ENTERED AT 13:38:56 ON 31 JUL 2008

SEA (POLONIUM OR PO OR LEAD OR PB) (A) 210 OR PB210 OR 210PB OR 2

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120
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    FILE ALUMINIUM
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     FILE ANTE
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2100 FILE AQUASCI
133 FILE AQUIRE
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  7
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     FILE CROPU
     FILE CSNB
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     FILE DDFB
 31
  9
     FILE DDFU
     FILE DISSABS
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     FILE DRUGB
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 11
     FILE DRUGU
 17
     FILE ELCOM
  7
     FILE EMA
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     FILE EMBAL
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5527 FILE ENERGY
380 FILE ENVIROENG
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125 FILE EPFULL 207 FILE ESBIOBASE

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    FILE FROSTI
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    FILE FSTA
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     FILE GBFULL
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     FILE GENBANK
    FILE GEOREF
3560
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95 FILE IFIPAT
5384
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 84 FILE INPADOCDB
 62 FILE INPAFAMDB
    FILE INSPEC
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147 FILE INSPHYS
    FILE IPA
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  3 FILE ITRD
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    FILE JAPIO
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    FILE KOREAPAT
     FILE LIFESCI
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 2
     FILE MATBUS
     FILE MECHENG
148
925
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118
     FILE METADEX
    FILE NAPRALERT
 1
    FILE NLDB
 43
941 FILE NTIS
     FILE OCEAN
756
 3 FILE PAPERCHEM2
1618 FILE PASCAL
 4 FILE PATDPA
    FILE PATDPAFULL
 30
    FILE PCI
 9
236
     FILE PCTFULL
     FILE PHIN
  1
  4
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     FILE PROMT
     FILE RAPRA
  2
     FILE RSWB
  3
     FILE RUSSIAPAT
  2
3578
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 38 FILE SOLIDSTATE
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    FILE TOXCENTER
4517
    FILE TRIBO
 1
102
     FILE TULSA
 22
     FILE TULSA2
 46
    FILE UFORDAT
117 FILE ULIDAT
1110
    FILE USPATFULL
333 FILE USPATOLD
145 FILE USPAT2
1271 FILE WATER
124 FILE WPIDS
124 FILE WPINDEX
  7 FILE WSCA
  QUE ABB=ON (POLONIUM OR PO OR LEAD OR PB) (A) 210 OR PB210 OR
   210PB OR 210PO OR PO210
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L36

D RANK

FILE 'STNGUIDE' ENTERED AT 13:40:57 ON 31 JUL 2008

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L38
          17263 SEA ABB=ON (LEAD OR PB)(A) 210 OR 210PB OR PB210
L39
        2217612 SEA ABB=ON FILM#
          68820 SEA ABB=ON RADON OR 222RADON OR RADON222
L40
         98370 SEA ABB=ON ALPHA(2A)(SOURCE OR EMIT? OR PARTICLE#)
L41
         47252 SEA ABB=ON POLYCARBONATE# OR POLY CARBONATE# 190082 SEA ABB=ON HYDROXIDE#
L42
L43
L44
         226644 SEA ABB=ON SEAL?
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L45
                OR L44)
L46
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            694 SEA ABB=ON L37 AND L38 AND L40
L47
            252 SEA ABB=ON L37 AND L38 AND L41 2 SEA ABB=ON L37 AND L38 AND L42
L48
L49
L50
             24 SEA ABB=ON L37 AND L38 AND L43
             21 SEA ABB=ON L37 AND L38 AND L44
L51
L52
             57 SEA ABB=ON L37 AND L38 AND (L39 OR L42 OR L43 OR L44)
             35 DUP REM L52 (22 DUPLICATES REMOVED)
L53
                     ANSWER '1' FROM FILE PASCAL
                     ANSWERS '2-11' FROM FILE BIOSIS
                     ANSWERS '12-29' FROM FILE ENERGY
                     ANSWERS '30-31' FROM FILE EMBASE
                     ANSWERS '32-34' FROM FILE COMPENDEX
                     ANSWER '35' FROM FILE SCISEARCH
              9 SEA ABB=ON L37 AND L38 AND L39 AND ((L40 OR L41 OR L43 OR
L54
                L44))
L55
             94 SEA ABB=ON L37 AND L38 AND L40 AND ((L41 OR L43 OR L44))
              6 SEA ABB=ON L37 AND L38 AND L43 AND ((L40 OR L41 OR L44))
L56
              4 SEA ABB=ON L37 AND L38 AND L44 AND ((L40 OR L41))
L57
           2389 SEA ABB=ON L37(2A) L38
L58
            501 SEA ABB=ON L58 AND (L40 OR L41)
            36 SEA ABB=ON L58 AND L40 AND L41
L60
            907 SEA ABB=ON L40(5A) COLLECT?
L61
              0 SEA ABB=ON L58 AND L61 AND L41
L62
              1 SEA ABB=ON L58 AND L61
L63
                D SCAN
L64
             29 DUP REM L60 (7 DUPLICATES REMOVED)
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                     ANSWERS '3-18' FROM FILE ENERGY
                     ANSWERS '19-26' FROM FILE INSPEC
                     ANSWER '27' FROM FILE EMBASE
                     ANSWERS '28-29' FROM FILE SCISEARCH
L65
             36 SEA ABB=ON L60 NOT (L49 OR L54 OR L56 OR L57 OR L63)
                D SCAN L64
L66
           1616 SEA ABB=ON RANDOM PULS?
              3 SEA ABB=ON L60 AND L66
L67
                D QUE
L68
              5 SEA ABB=ON L58 AND (L40 OR L41) AND L66
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L69
                D SCAN
                D TRIAL
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E K08-A+ALL/MC
L70
             68 SEA ABB=ON (POLONIUM/BI, ABEX OR PO/BI, ABEX) (A) 210/BI, ABEX OR
                210PO/BI, ABEX OR PO210/BI, ABEX OR POLONIUM210/BI, ABEX OR
                210POLONIUM/BI, ABEX
             68 SEA ABB=ON (LEAD/BI, ABEX OR PB/BI, ABEX) (A) 210/BI, ABEX OR
L71
                210PB/BI, ABEX OR PB210/BI, ABEX OR 210LEAD/BI, ABEX OR LEAD210/BI
                , ABEX
L72
             11 SEA ABB=ON L70 AND L71
                D SCAN
         974355 SEA ABB=ON FILM#/BI,ABEX
L73
         209521 SEA ABB=ON ALPHA/BI,ABEX
L74
         66573 SEA ABB=ON POLYCARBONATE#/BI,ABEX OR POLY CARBONATE#/BI,ABEX 146392 SEA ABB=ON HYDROXIDE#/BI,ABEX
L75
L76
L77
         722352 SEA ABB=ON SEAL?/BI,ABEX
           1259 SEA ABB=ON RANDOM PULS?/BI,ABEX
L78
L79
              9 SEA ABB=ON L70 AND L71 AND (L73 OR L74 OR L75 OR L76 OR L77
                OR L78)
           2652 SEA ABB=ON ALPHA/BI, ABEX(2A) (SOURCE/BI, ABEX OR EMIT?/BI, ABEX
L80
                OR PARTICLE#/BI, ABEX)
              4 SEA ABB=ON L70 AND L71 AND (L73 OR L75 OR L76 OR L77 OR L78
L81
                OR L80)
     FILE 'STNGUIDE' ENTERED AT 14:07:22 ON 31 JUL 2008
     FILE 'CAPLUS' ENTERED AT 14:09:11 ON 31 JUL 2008
                D QUE L9
                D QUE L16
                D OUE L25
                D QUE L26
                D QUE L32
                D QUE L35
             14 SEA ABB=ON (L9 OR L16 OR L25 OR L26 OR L32 OR L35)
L82
     FILE 'WPIX' ENTERED AT 14:09:14 ON 31 JUL 2008
                D OUE L81
     FILE 'PASCAL, BIOSIS, GEOREF, ENERGY, DISSABS, CONFSCI, INSPEC, EMBASE,
     COMPENDEX, SCISEARCH' ENTERED AT 14:09:14 ON 31 JUL 2008
                D QUE L49
                D QUE L54
                D QUE L56
                D QUE L57
                D QUE L63
                D OUE L68
L83
             23 SEA ABB=ON (L49 OR L54 OR L56 OR L57 OR L63 OR L68)
     FILE 'CAPLUS, WPIX, BIOSIS, ENERGY, INSPEC, EMBASE, COMPENDEX, SCISEARCH'
     ENTERED AT 14:09:22 ON 31 JUL 2008
L84
             31 DUP REM L82 L81 L83 (10 DUPLICATES REMOVED)
                      ANSWERS '1-14' FROM FILE CAPLUS
                      ANSWERS '15-16' FROM FILE WPIX
                     ANSWERS '17-20' FROM FILE BIOSIS
                     ANSWERS '21-28' FROM FILE ENERGY
                      ANSWER '29' FROM FILE EMBASE
                      ANSWER '30' FROM FILE COMPENDEX
                     ANSWER '31' FROM FILE SCISEARCH
                D IBIB AB HITIND 1-14
                D IALL ABEX TECH 15-16
                D IALL 17-31
```

FILE 'HOME' ENTERED AT 14:09:55 ON 31 JUL 2008